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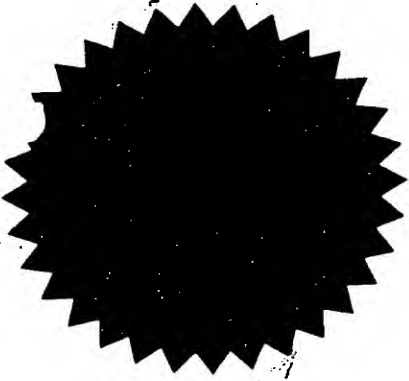
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חוק הפטנטים, תשכ"ז - 1967  
PATENTS LAW, 5727-1967

ב ק ש ה ל פ ט נ ט  
Application For Patent

מספר: Number	121512
תאריך: Date	14-07-1997
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אני, (שם המבקש, מענו ולגבי גוף מאוגד - מקום התאגדותו)  
I, (Name and address of applicant, and in case of body corporate-place of incorporation)

מוסד הטכניון למחקר ופיתוח בע"מ, חברה ישראלית, מבית הסנט, פארק גוטווירט, קרית הטכניון,  
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(בעברית)  
(Hebrew)

Microelectronic components, their fabrication and electronic networks  
(English)

comprising them hereby apply for a patent to be granted to me in respect thereof. מבקש בזאת כי ינתן לי עליה פטנט

* בקשת חלוקה - Application of Division		* בקשת פטנט מוסף - Appl. for Patent of Addition		* דרישת דין קדימה Priority Claim	
מבקשת פטנט from application		* לבקשה/לפטנט to Patent/Appl.		מספר/סימן Number/Mark	תאריך Date
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REINHOLD COHN AND PARTNERS Patent Attorneys P.O.B. 4060, Tel-Aviv C. 106906					
חתימת המבקש Signature of Applicant				היום 14 בחודש July 1997 שנת of the year of This	
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המכילות אותן

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Microelectronic components, their fabrication and electronic networks  
comprising them

Technion Research and  
Development Foundation Ltd.

מוסד הטכניון למחקר ופתוח בע"מ

The inventors:

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C. 106906

## MICROELECTRONIC COMPONENTS, THEIR FABRICATION AND ELECTRONIC NETWORKS COMPRISING THEM

### FIELD OF THE INVENTION

The present invention is generally in the field of microelectronics and concerns components for use in microelectronics, manner of fabrication of such components as well as networks, particularly microelectronic  
5 networks, making use of these components.

### PRIOR ART

Prior art believed to be relevant as background of the invention as well as to manufacture or experimental techniques described herein is  
10 listed below:

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35 Acknowledgement of these references will be made by indicating these numbers from the above list.

## BACKGROUND OF THE INVENTION

40 The miniaturization of microelectronics and logics, at current technology, is approaching its practical and theoretical limits. Various design and operational considerations such as heat removal, heterogeneity, connectivity as well as present photolithographic techniques restrict the practical size of minimal feature is present, semiconductor-based electronic

components to about 0.25–0.3  $\mu\text{m}$ . It is clear that further miniaturization of electronic components must involve new approaches and concepts for the fabrication of the electronic components and logic circuits.

5 Nanometer scale electronics needs to consider two fundamental issues: operating principles of the corresponding electronic components and schemes to fabricate such components and their integration into useful circuits.

A number of operation principles have been suggested based on charging effects<sup>(1-6)</sup> which become increasingly prominent as the device dimensions diminish. The construction of nanoscale circuits cannot be implemented by existing microelectronics technology. In particular, inter element wiring and electrical interfacing to the macroscopic world become increasingly problematic. Molecular recognition processes and self-assembly of molecules into supramolecular assemblies may be used for the construction of complex structure<sup>(5)</sup>. However, integrating electronic materials with these structures, or providing them with electronic functionality, has not yet been attained.

20 Nucleic acids possess self-assembly properties which can be used to form networks of nucleic acid fibers<sup>(27-30)</sup>. DNA has already been employed as an organizer of nano structures in the assembly of colloidal particles into macroscopic crystal-like aggregates<sup>(14,15)</sup> and in dictating the shape of semiconductor nano particle assemblies<sup>(16,17)</sup>.

## GLOSSARY

25 In the following, use will be made with some terms, which terms and their meaning are as follows:

*Nucleic acid fiber* (to be referred to at times as "*fiber*") – A sequence of nucleotides, which may be ribonucleotides, deoxyribo-nucleotides, other ribo nucleotide derivatives, a variety of synthetic, i.e. non naturally occurring nucleotides, as well as any combination of the above. The nucleic acid fiber may be single-stranded, double-stranded or multi-stranded.

*Binding* – A term to refer collectively to all types of interactions which bind together two or more molecules, substances, particles, supramolecular structure, etc., or brings to binding of these to a solid substrate. The binding may be covalent or non-covalent binding (non-covalent binding may include one or more of ionic interaction, hydrophobic interaction, Van der Waals interaction, chemical sorption, etc.). Derivation of this term (i.e. "bind", "bound", etc.) will have meanings commensurate to that assigned to "binding", depending on the syntax.

10 *Functionalized fiber* – A nucleic acid fiber carrying a substance or particles which impart electric or electronic properties to the fiber. Such substances or particles may be bound to the fiber by a variety of interactions (e.g. may be chemically deposited on the fiber, may be complexed thereon by a variety of chemical interactions, may be associated with the fiber by  
15 electrostatic or hydrophobic interactions, etc.). The substances or particles may be bound to the fiber based on the general chemical properties of the fiber or may be bound to the fiber in a sequence specific manner. Examples of such substance or particles are: metal e.g. which gives rise to a conductive fiber; a variety of semi-conductive materials which can form conductors  
20 (or resistors), electronic p/n junctions, and others; colloid particles which can be associated with the fiber to form functional logic junctions between fibers; etc. Depending on the type of bound substances or particles, the electric or electronic properties imparted to the fiber may be conduction, insulation, gating, switching, electrical amplification, etc.

25

*Junction* – A point of connection of two or more fibers to one another. Examples of junctions can be seen in Fig. 2 (please refer to the description below relating to this figure).

30 *Network* – A geometrical one, two or three-dimensional structure made of functionalized fibers creating a network and other compounds such as colloid particles, proteins, other macromolecules and supramolecular

complexes, etc. The term "*network*" may refer to the geometrical arrangement of nucleic acid fibers which comprises the fibers and junctions between them. This term may also be used, at times, collectively to denote also a *functionalized network* (see below).

5

***Functionalized network*** - a network comprising at least one functionalized fiber. The functionalized network has properties defined among others, by the type of connectivities and *junctions* between different fibers, by the pattern of the *interface components* (see below) and by the nature of the deposited or complexed substances or particles.

10

***Interface component*** - A conducting substrate, which may be made of metal or of any other conducting material or coated by metal or such conducting material, which serves for connection of said network or to external electronic or electric components or circuitry, thus serving as an input/output (I/O) interface with an external component or circuitry. The interface components are linked to the network on the one hand and are electrically linked to an external component or circuitry, on the other hand. The interface component, may at times also refer to herein as "*electrode*".

20

***External circuitry, external component*** - An electronic or electric circuitry or an electronic or electric component, respectively, situated electrically external to the network and typically comprises prior art electric or electronic components, including standard solid-state microelectronic components.

25

***Linker*** - An agent (molecule, complex of molecules, supermolecular structure, macromolecule, aggregate, colloid particle, molecular clusters, etc.) that acts in providing a physical link between the network and the interface component. The linkers may have chemical groups for covalent or non covalent binding, (e.g. complexation or sorption, etc.) to the interface

30

components, on the one hand, and to the nucleic acid fiber of the network on the other hand. Examples of a linker are: a nucleic acid binding protein; synthetic molecules with a binding ability to a specific nucleic acid sequence; a short, single or multiple stranded nucleic acid sequences (e.g. an oligonucleotide), e.g. having a "sticky end" and being modified at its other end, to allow it binding to the interface component; etc.

**Complexing agent** - An agent which is used for binding of components of the network to one another. The binding may be covalent or non covalent. Examples of complexing agents are: proteins with a specific binding ability to a nucleic acid sequence; oligonucleotides; synthetic molecules which can bind to two components; etc. An example of use of a complexing agent is in linking of an end of a nucleic acid fiber to a colloid particle or linking of the two nucleic acid fibers to one another, for the purpose of creating a junction.

**Wire** - In the context of the invention - a functionalized fiber with bound substance or particles which give rise to electric conductivity along the fiber. The wires are conducting components which may interconnect two or more sites of the network, connect the network with an electrode, connect between two networks, etc. A wire may extend an entire length of a functionalized fiber or may extend only part of the length of a functionalized fiber, with other parts serving as base for various types of electronic components, e.g. a diode, an electric switch, etc.

25

**Insulator** - A fragment or component which acts as a barrier for electric conduction.

**Switch** - A two or multi terminal component where the conductance between any pair of terminals can be turned on or off by a control signal including, for example, the potential at another terminal, light, pressure, chemical reaction, stress, etc.

30

*Electronic component* – Any component in a functionalized network other than a wire or a simple junction between wires (a simple junction being a junction having the purpose of only providing a link between fibers or functionalized fibers).

5

*Electronic functionality* (or *electronically functional*) – A property of a component in a functionalized network which renders it to serve as an electronic component.

## 10 SUMMARY OF THE INVENTION

The present invention makes use of the molecular recognition properties and self-assembly processes of nucleic acid sequences and other components. These features are used to prepare nucleic acid fiber-based networks with a geometry defined by the type of interconnectivity between  
15 the fibers. A network with electronically functional properties (functionalized network) is obtained by binding substances or particles to the nucleic acid fibers. The functionalized network may include conductors, switches, diodes, transistors, capacitors, resistors, etc.

The present invention provides, by a first of its aspects, a  
20 microelectronic circuit comprising a network of at least one electric or electronic component, the network being connected to at least one, preferably at least two input/output interface components; characterized in that

the network has a geometry defined by at least one nucleic acid fiber;  
25 each of said interface components is connected to at least one fiber; and in that

said at least one fiber has one or more substances or particles bound thereto or complexed therewith such that at least one electric or electronic component is formed with properties based on electric charge transport  
30 characteristics of said one or more substance or particles, and typically also on its location and connectivity within the network.

The microelectronic circuit comprises at least one conductor and at least one microelectronic component. The microelectronic network has typically the following characteristics:

- 5 (i) the conductor is a wire constructed on at least one first portion of the at least one nucleic acid fiber which is electrically functionalized by having one or more electrically conductive substances or particles bound thereto such that electric current can flow along said at least one first portion of the nucleic acid fiber;
- 10 (ii) the microelectronic component being electrically connected to said at least one wire and being constructed either on at least one second portion of a nucleic acid fiber, other than said at least one first portion, by having one or more substances or particles bound thereto rendering said at least one second portion to have electronic functionality (i.e. serving as an electronic component of the network) or being constructed by a particle situated at a  
15 junction between two or more nucleic acid fibers rendering said junction to assume an electronic functionality, said electronic functionality being based on electric charge transport characteristics of said one or more substance or particles (i.e. serving as an  
20 electronic component of the network);
- (iii) each of said input/output interface component is connected to at least one functionalized nucleic acid fiber or at least one of said microelectronic components in a manner allowing electric  
25 conductivity between said interface component and said functionalized fiber or such microelectronic component, respectively.

The skeleton of the network of the invention comprises nucleic acid fibers which are assembled to form a network on the basis of their sequence specific hybridization with other fibers or specific binding to other  
30 components. In this manner networks with practically infinite variety of geometries can be formed.

Substances or particles may be bound to the nucleic acid fibers based on the general (overall) chemical properties of the fibers. This will typically yield a substantially homogeneous deposition of the substance or particles along the nucleic acid fibers. A specific embodiment of such a homogeneous binding of substances or particles is the formation of an electrical conducting wire, e.g. where the conducting substance is metal, such as described below. Alternatively, substances or particles may also be bound to the fibers in a sequence or domain-specific manner in different portions of the fibers, namely in a manner which depends on the sequence of nucleotides at given portions of the nucleic acid fibers.

Sequence or domain-specific deposition of substances at different fiber portions may be performed in a number of different ways. For example, an oligonucleotide *a priori* bound to a certain electronically functional substance or particle, may be made to bind onto a fiber with a complementary sequence. Similarly, it is possible also to bind different types of substances or particles, in a sequence or domain-specific manner, also to a multi-stranded (e.g. double-stranded) nucleic acid fiber. This may be achieved, for example, by the use of a sequence-specific complexing agent which identifies and binds to a specific site of a double-stranded nucleic acid fiber. The complexing agent may be an oligonucleotide, forming with a double-stranded fiber, a triple-stranded structure; a protein recognizing a specific double-stranded domain; and many others.

By sequence or domain-specific binding, different types of substances may be bound to different portions of a given fiber or network of fibers.

Particles, e.g. colloid particles or polymers, may be made to bind to one or more fibers, typically by the use of complexing agents. Depositions of such particles may be utilized for the formation of electronic components, e.g. a single-electron transistor (SET).

The geometry of the network is defined by the nucleic acid fibers. In formation of the network the chemical complementary and molecular recognition properties are utilized by employing self-assembly processes.

The nucleic acid fibers are assembled to form the network by specific hybridization of complementary sequences to form local or global minimum-energy structures. This may be used for the formation of various junctions (e.g. T- or X- junctions, as exemplified in Fig. 2 and others).  
5 Specific molecular recognition between nucleic acid fibers and linkers or complexing agents (the linkers or complexing agents may be oligonucleotides or a variety of other molecules, macromolecules, supramolecular assemblies or particles) may be used to link the nucleic acid fibers to interface components or to other particles situated at junctions between  
10 fibers.

The invention also provides, by another of its aspects, a method for fabricating a microelectronic network connected to interface components comprising

- (a) providing an arrangement consisting of at least one (preferably at  
15 least two) interface component;
- (b) immobilizing a linker on said interface component capable of binding to a nucleic acid sequence,
- (c) contacting said interface component with at least one nucleic acid fiber with a sequence capable of binding to the immobilized linker, and  
20 providing conditions for binding of said sequence to said deposited agent,
- (d) functionalizing the at least one fiber by binding thereto, depositing thereon or complexing therewith substance or particles such as to form electric or electronic components and said at least one fiber, the function of said electric or electronic component being based on charge  
25 transport properties of said substance or particles.

It should be noted that the order of steps in the above method is not material and may be changed. For example, step (c) may precede step (b), or the functionalizing step (d) may precede step (c).

The network may at times be formed at once by mixing all  
30 components in a medium and then allowing the components to self-assemble in a specific manner, based on the pre-designed properties of the various components. Nucleic acid fibers may be designed to have specific sequences

to allow their hybridization to complementary sequences in other strands. Similarly, particles may be formed with specific, sequence or domain-recognizing complexing agent bound thereto, to allow them to bind to nucleic acid fibers in a sequence or domain-specific manner. For example, 5 colloid particles with three different oligonucleotides bound thereto can be formed which will then bind to ends of three different fibers, to form a junction between the three fibers containing a colloid particle. Similarly, in order to ensure binding of the network to the interface components in a specific manner, sequence- or domain- recognizing linkers are immobilized 10 on the interface components which is brought into contact with the assembling network.

At other times the network may be formed in a sequential manner, e.g. first forming a first sub-network structure comprising part of the components of the complete network and then the missing components 15 (e.g. fibers, particles, etc.) may then be sequentially added until the network is completed. Sub-network structures may, for example, be particles with several fibers connected thereto, branched - fiber structures, etc. The gradual assembly may also be based on the self-assembly properties of the nucleic acid fibers and of complexing agents and linkers which bind to the 20 fibers in a sequence or domain-specific manner. Furthermore, it is also possible, particularly in the case of networks with a complex structure, to first prepare a plurality of sub-network structures and then combine them for the formation of the complete network.

As will obviously be appreciated by the artisan, the formation of 25 the network may be aided by agitating the medium where the network is formed, by providing directional streams of fluid to orient the fibers to connect to a downstream component at their other end, by applying other biasing measures, etc. In forming wires of the inventions, electric potential between the two ends of the fiber on which the wire is formed, may enhance 30 and provide directional deposition of the metal.

The nucleic acid fibers, which may be *a priori* single, double or multiple stranded, may be formed and replicated by a variety of methods

including recombinant DNA methods involving production and reproduction of nucleic acid fibers by "*engineered*" cells, e.g. microorganisms; alternatively, the fibers may be produced synthetically, e.g. by synthesis of strands and then combining them into larger fibers. The fibers may be formed by  
5 a variety of amplification techniques, e.g. polymerase chain reaction (PCR); etc.

The invention also provides by another of its aspects, electronic components useful in said functionalized network, but which may have a further utility by their own right. Examples of such electronic components  
10 are: switch, diode, transistor, SET, field effect transistor (FET), wire, capacitor, resistor, etc.

A wire of the invention may be made to be very thin and may be used to advantage in application requiring thin wires, for example, as a gate in a semi-conductor FET for a very fast gating of such a transistor. The  
15 gating speed of a FET depends to a large extent on the width (referred to in the art at times as "*length*") of the gating wire in the FET. The wire in accordance with the invention may be made to be about two orders of magnitudes smaller than the width of prior art gate wires of FETs, and accordingly, fast modulation, faster than hitherto possible, can be obtained  
20 in a FET using a wire of the invention. A FET comprising a wire of the invention as its gate is also an aspect of the invention.

The invention will now be illustrated by the following detailed description and subsequent examples, with occasional reference made to the annexed drawings. It should be appreciated by the artisan, that the invention  
25 is not limited to the specifically described embodiments but rather applies to the full scope of the invention as defined above, namely, to the formation of a network, and components in the network, by employing self-assembly properties of nucleic acid fibers and by deposition or complexation of substances or particles onto or to the fibers using molecular recognition  
30 driven, self assembly process, to render the fibers or junctions between the fibers electronically functional.

## BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1A shows a matrix of interface components and linkers for binding to the network of the invention on the one hand and connect to external electronic circuitry or components on the other hand.

5 Fig. 1B illustrates an embodiment of immobilization of oligonucleotide linkers onto the interface components.

Fig. 2 shows examples of junctions between nucleic acid fibers (six examples are shown in Figs. 2A-2F, which are explained in the text below).

10 Fig. 3A is a presentation of the manner of forming and functionalizing a nucleic acid fiber into a wire, with the conducting material being silver.

Fig. 3B shows a possible current-voltage relationship of a wire formed as illustrated in Fig. 3A, which is dependent on the scanning direction (presented by arrows on the curves).

15 Fig. 4 is a representation of the manner of forming a wire, where the electronic material is polyphenylene vinylene (PPV).

Fig. 5 shows a number of examples of functionalized fibers consisting of a p/n junction (Figs. 5A-5C), a graded p/n junction (Fig. 5D) and a bipolar n-p-n transistor (Fig. 5E).

20 Fig. 6 is a presentation of the manner of forming a single electron transistor (SET) by one embodiment of the invention.

Fig. 7 is an illustration of a molecular switch in accordance with an embodiment of the invention.

25 Fig. 8 is an illustration of a FET in accordance with an embodiment of the invention, shown in a planar view (Fig. 8A) and in a cross-sectional view (Fig. 8B).

Fig. 9 shows the scheme for synthesizing an oligonucleotide, as described in Example 1(A).

Fig. 10 shows a fluorescently labeled  $\lambda$ -DNA stretched between two gold electrodes (dark strips) 16  $\mu\text{m}$  apart.

**Fig. 11** shows atomic force microscope (AFM) images of a silver wire connecting two gold electrodes 12  $\mu\text{m}$  apart (a) 1.5  $\mu\text{m}$  and (b) 0.5  $\mu\text{m}$  field sizes.

**Fig. 12A** is two terminal I-V curves of the silver wire prepared according to Example 9. The arrows indicate the voltage scan direction. The solid-line curves are repeated scans and demonstrate the stability of the samples. Note the different asymmetry in the I-V curves corresponding to the two scanning directions.

**Fig. 12B** shows the I-V curves of a different silver wire in which the silver growth was more extensive than in Fig. 12A. The more extensive silver growth resulted in a smaller current plateau, on the order of 0.5V, and a lower resistance (13M $\Omega$  vs. 30 M $\Omega$  in Fig. 12B). By driving large currents through the wire, the plateau has been eliminated to give an ohmic behavior (dashed line), over the whole measurement range.

## DETAILED DESCRIPTION OF THE INVENTION

The formation of a network typically begins by providing a matrix of interface components, which provide the I/O interface between the network and an external circuitry or external component. Illustration of one embodiment of such an interface component matrix 100 is shown in Fig. 1A. Each of the interface components 102 is typically a metal electrode having a network-connecting pad 104, and an external circuitry connecting pad 106 linked by a connector portion 108.

As a preparatory step for the formation of the network, the connecting pads are treated to allow binding thereto of linkers 110, e.g. derivatized oligonucleotides. Examples of the manner of treatments are described below. (One embodiment of the manner of immobilizing a linker oligonucleotide onto an interface component is seen in Fig. 1B and described below). Pre-prepared linkers 110 can then be attached to the pad 104, typically a different linker to each of the pads. The linkers 110, shown in a schematical manner in the enlargement at the right of the central

portion 112 of the matrix, may be immobilized onto pads 104, for example, by jet printing, e.g. in the manner described below in the Examples. In this manner, a different linker 110 may be attached to each of pads 104. Each of linkers 110 may have a selective binding ability to a different specific  
5 nucleic acid sequence, this feature being represented by the different shapes at the end of the linkers.

A functionalized nucleic acid network comprising functionalized fibers including wires (as herein defined and described, by way of example, below) and a variety of electronically functional components formed on the  
10 nucleic acid fibers or at junctions between the fibers, can then be contacted with the matrix; the specific binding of nucleic acid sequences in the network's fibers to the linkers immobilized onto the interface components results in a specific connectivity pattern of the network to the interface components. Alternatively, nucleic acid fibers may first be made to  
15 specifically bind to the linkers and the functionalization of the networks, i.e. the formation of electronically functional components, the formation of wires, etc., may then be carried out in the formed nucleic acid-based network. Another alternative would be to first anchor several sub-network structures to the matrix of interface components and at a subsequent step, or  
20 simultaneously, allow the sub-network to bind one another to form a complete network. Examples of sub-network structures include several nucleic acid fibers connected together forming part of the fully-fledged network; colloid particles with several nucleic acid fibers attached thereto; etc. Here again, the fibers may be *a priori* electronically functionalized  
25 (wholly or partially) or the functionalization may be carried out after formation of the nucleic acid fiber network.

Fig. 1B is a schematic representation of one embodiment for immobilizing an oligonucleotide onto an interface component. Biotin molecules 120 and 122 are bound, one to an oligonucleotide 124 and the  
30 other to a sulfur containing moiety 126. Biotin molecule 122 is immobilized onto interface component 128 through the sulfur containing moiety 126

and then when streptavidin molecules 130 are introduced into the medium, they yield the formation of a complexing agent 132, which is a supermolecular complex comprising biotin and streptavidin, which immobilizes oligonucleotide 124 onto interface component 128.

5            Important components in the network of the invention are junctions which serve a variety of functions. Several examples of junctions are shown in Fig. 2. Fig. 2A depicts a junction 200, formed between two single-stranded nucleic acid fibers 202 and 204. The junction, in this specific example is formed by hybridization of a terminal end sequence 206  
10 in fiber 204 and a complementary sequence 208 in fiber 202. This junction may serve as a T-type junction between nucleic acid fibers which can then be transformed into functionalized connecting junctions by depositing an electrically conducting substance on fibers 202 and 204 and junctions, e.g. in the manner to be described below.

15            Another type of junction 210 is shown in Fig. 2B. This junction is formed by a complexing agent 212 which may be a colloid particle, a protein, another type of a macromolecule, a supramolecular structure, etc. In this case, the junction is formed between one single-stranded fiber 214 and another single-stranded fiber 216. However, it should be appreciated  
20 that such a junction may also be formed between two double-stranded fibers. The complexing agent may be bound to the fibers by a variety of means. The binding may be covalent or non-covalent. Examples of non-covalent binding are ionic interactions, hydrophobic interactions, by means of Van der Waals forces, etc. The complexing agent may also be a complex  
25 molecular structure by itself, e.g. it may be formed by two or more molecules or macromolecules with a binding affinity to one another of which at least one is bound to each of fibers 216 and 214. Examples of such complex molecular structures are a large variety of molecules which can bind to one another, such as for example: antibody-antigen, ligand-  
30 receptor, biotin-avidin, and many others.

A specific example of such a complex molecular structure is shown in Fig. 1B (above). Although this complex molecular structure, which is a biotin-avidin complex, is shown in its role as immobilizing oligonucleotide onto an electrode, the same complex molecular structure  
5 may also be used for forming a junction between nucleic acid fibers (i.e. each fiber will be modified by binding a biotin derivative and then the two avidin moieties may be complexed by one streptavidin).

A further example of a junction 220 is shown in Fig. 2C. In this case, a complexing agent 222 binds together two fibers 224 and 226. The  
10 complexing agent and the manner of its interaction with nucleic acid fibers may be similar to complexing agent 212 in Fig. 2B.

Two other examples of junctions are shown in Figs. 2D and 2E. These junctions 230 and 240 are formed between three and four fibers, respectively, each of which is double-stranded. Such junctions may be  
15 formed by hybridization or enzymatic synthesis, as described below in the Examples.

Another type of junction 250 is shown in Fig. 2F. In this case, a complexing agent 252, e.g. a colloid particle, links together a plurality of fibers, four 253-256 in this specific embodiment, each of which is bound  
20 at its terminal to the complexing agent 252. The binding of each of fibers 253-256 to particle 252 may be by means of direct association, by the use of mediators such as specific binding proteins, in any of the manners described in connection with Fig. 2B and 2C, etc.

In the following, some examples of fabrication of wires and  
25 electronic components and functionalized networks of the invention will be described. It should be understood that these are exemplary embodiments only and various modifications of the described embodiment are possible, all being within the scope of the invention as defined herein.

Reference is now being made to Fig. 3, showing the manner of  
30 formation of a wire in accordance with the invention between two electrodes 300, typically made of, or coated by metals such as gold,

platinum, silver, etc. Electrodes 300, which may be first treated in a manner to facilitate subsequent binding of the linker, are wetted separately with a solution containing either linker molecules 302 or 304, each consisting of a single-stranded oligonucleotide ("*Oligo A*" and "*Oligo B*", respectively),  
5 derivatized by a disulfide group. When these linkers are deposited on electrodes 300, under appropriate conditions, the disulfide group bind to the electrodes 300, to form linkers 306 and 308, respectively (step (a)). Electrodes 300 are then wetted with a nucleic acid fiber solution, e.g. a DNA double-stranded fiber, 310, having sticky ends, complementary to the  
10 sequences of the oligonucleotides in linkers 306 and 308. Electrodes 300 are spaced from one another at a distance about equal to the length of the nucleic acid fiber 310, whereby each end of nucleic acid fiber 310 is connected to its complementary oligonucleotide in one of linkers 306 and 308 to form a bridge 312 between the two electrodes 300 (step (b)). By  
15 controlling the concentration of oligonucleotide 310 in the medium, the number of such bridges formed between the electrodes can be controlled. Following hybridization the binding of the linkers to the nucleic acid fibers may be strengthened by covalent binding of the two to one another by ligation of the nicks.

20 At times, particularly where fiber 310 is long and thus it is not practical to ensure its hybridization at both ends merely by relying on diffusion, the strand 310 may be made to connect to one electrode and then, by a directional stream of fluid from the first electrode to the second, the nucleic acid fiber is made to extend so that its end reaches the second  
25 electrode.

It is also clear that in order to avoid folding of the nucleic acid fibers and to ensure proper binding, appropriate solutions may need to be at times selected.

The functionalization step of the fiber, for the purpose of  
30 constructing a metal wire, begins, according to the specifically illustrated embodiment, by an ion exchange step involving exposure of the fiber to a

solution comprising silver ions ( $\text{Ag}^+$ ) under alkaline conditions, whereby the silver ions replace the sodium ions or other ions normally associated with the fiber and complex with the negatively charged fibers (step (c)). This gives rise to a nucleic acid fiber 314 loaded with silver ions 316. It should  
5 be noted, that rather than silver ions, a wide variety of other metal ions can be used, including for example, cobalt, copper, nickel, iron, gold, etc. Furthermore, metal aggregates, complexes or clusters, e.g. colloidal gold, colloidal silver, etc., may also be deposited on the nucleic acid fiber via a variety of different interactions. The ion-exchange step typically involves  
10 rinsing of the fibers with deionized water, then drying the fibers and then soaking them in a solution of the metal ions or metal aggregates.

At a next step (step (d)), the fiber is exposed to a reducing agent, e.g. hydroquinone, which yields a reduction of the metal ions *in situ* into metallic silver. The metallic deposit, e.g. metallic silver, is formed at a  
15 number of nucleation sites 318. After rinsing with deionized water and drying, the fibers with nucleation sites 318 are contacted with a reagent solution comprising metal ions and a reducing agent, e.g. hydroquinone, under acidic conditions. Under these conditions, the ions are converted to metallic metal only at the nucleation site and consequently the nucleation  
20 centers grow to form a conductive wire 320 (step (e)).

The so formed wire 320 may be subjected to a variety of post fabrication treatments, which may include, for example, thermal treatment intended to increase the wire's thickness and homogeneity; passivation treatment for the purpose of forming an electrically insulating layer around  
25 the wire, e.g. by exposure to alkane thiol; electrochemical or photochemical growth of the wire by polymers on the wires; etc.

Fig. 3B illustrates a typical current-voltage relationship of a wire formed by the procedure illustrated in Fig. 3A. As can be seen, the curves are non linear and are asymmetric with respect to zero bias. The shapes of  
30 the curves depend on the fold of scan direction as indicated by the arrows in Fig. 3A. Approaching zero voltage from a large positive or negative bias, the current vanishes almost linearly with the voltage. A zero current plateau

then develops with very large differential resistance. At a higher bias, the wire turns conductive again with a different channel resistance, lower than in the original bias polarity. This history-dependent current-voltage relationship, may render the wire as a logic or memory component.

5           Reference is now being made to Fig. 4, showing the manner of formation of a wire in accordance with another embodiment of the invention. However, rather than metal, as in the embodiment of Fig. 3, in this embodiment, the deposited material is PPV (poly-phenylene vinylene). Electrodes 400, may be the same as electrodes 300 shown in Fig. 3. The  
10 first two steps of the method (steps (a) and (b)), are substantially identical to the corresponding steps in Fig. 3 (identical components have been given a reference numeral with the same last two digits as the corresponding components in Fig. 3: e.g. 402 is the same as 302, 404 as 304, etc.). The formed bridge 412 may be strengthened, similarly as above, by covalent  
15 binding of fiber 410 to linkers 406 and 408 to yield a complete fiber 414 connecting the two electrodes (step (c)).

A solution comprising pre-PPV<sup>(26)</sup> molecules 416 is then contacted with fiber 414 and by the virtue of being positively charged, pre-PPV 414 becomes complexed with the negatively charged DNA bridge 414  
20 (step (d)). At a next step, the sample is rinsed, dried and finally heated in a vacuum, e.g. to a temperature of about 300°C, for about 6 hours, which leads to the release of tetrahydrothiophene groups and hydrochloric acid from each repeat unit, yielding a luminescent PPV component (step (e)).

In order to convert the PPV into a conductor, this polymer is  
25 doped with agents which either cause electron deficiency (holes) or give rise to extra electrons. Doping may be performed by many known methods e.g. exposure to H<sub>2</sub>SO<sub>4</sub> vapor, addition of halo-acid vapor (e.g. HCl, HBr), by the use of dodecyl benzene sulfonic acid, by the use of camphor sulfonic acid, or by other means. The extent of doping determines the conductivity  
30 of the wire.

Many other polymers may be used instead or in addition to PPV in accordance with the invention. This includes a variety of polymers with

positively charged side groups as well as polymers with positively charged groups in the backbone or polymers with recognition groups capable of binding to nucleic acid fibers. Another example of a polymer is polyaniline (PANI). These polymers include such which have either an electron deficiency (p-type polymers) or have electron surplus, (n-type polymers).  
5 In addition in a similar manner, *mutatis mutandis*, other types of conducting substances (n-type or p-type substances) may be bound to the fiber.

By the use of oligonucleotides bound to various substances which can impart electronic functionality, the properties of the electronic components assembled on a nucleic acid fiber can be precisely controlled. For  
10 example, two oligonucleotides, of which one has a sequence to allow hybridization to a specific portion of the fiber, and is bound to a p-type substance, and another has a sequence allowing hybridization to an adjacent portion of the fiber, bound to an n-type substance, (a polymer with an  
15 electron surplus) are made to bind to the fiber, and in this manner, an n/p junction can be formed, e.g. serving as a diode. Another example may be the formation of a p-n-p or an n-p-n type, bipolar transistor.

An example of some functional components which may be formed on a fiber are shown in Fig. 5. In Fig. 5A, a p/n junction is formed  
20 by a p-type substance 510 bound to one oligonucleotide 512, which is a poly C in this specific example, and an n-type substance 514 bound to another oligonucleotide sequence 516, a poly A in this specific example. The oligonucleotides bind to complementary sequences 518 and 520, respectively, on the DNA fiber 522 and after coupling, a p/n junction is  
25 formed. In Fig. 5A, the p/n junction is formed on a single-stranded segment of fiber 522. Similarly, such a junction may also be formed on a double-stranded fiber 524 (see Fig. 5B), e.g. by first removing a portion of one strand, e.g. by enzymatic digestion to expose adjacent segments 526 and 528 and then hybridization with the complementary p-type and n-type  
30 substance-carrying oligonucleotides 530 and 532, respectively.

The remaining portion of the fiber, may, for example, be treated in a manner to fabricate a wire, such as that described above with reference to Figs. 3 or 4, and accordingly a diode (a p/n junction) 534 is formed with conducting wires (C) 636 and 538 flanking the two ends of the diode (Fig. 5C). In order to ensure that the conductor substance is not deposited on the p/n junction portion, the materials constituting the junction may first be coupled and only then the remaining portion of the fiber may be treated in a manner described above to form a conducting wire.

Fig. 5D is a schematic representation of another structure which in this case consists of two conducting wires (C) formed on two peripheral fiber portions 540 and 542 which flank a portion of the fiber which was transformed into a graded p/n junction 544. This exemplified junction consists of portions which are heavily doped 546 and 548 (ppp and nnn) portions which are moderately doped 550 and 552 (pp and nn) and portions which are only slightly doped 554 and 556 (p and n).

Another functionalized fiber is shown in Fig. 5E. This wire has a T-type branch point 560 and by converting the portion of the fiber into an n-p-n bipolar junction 562 as illustrated (or a p-n-p junction) and then converting the remaining portions of the fibers 564, 566 and 568 into wires, a bipolar transistor is formed.

Specific depositions of various substances can also be achieved by means other than hybridization. Various molecules, e.g. proteins, which are capable of recognizing specific domains, even without the need to "open" the double or multi-stranded fiber, may be used for this purpose.

Fig. 6 shows a manner of construction of a single electron transistor (SET). A colloid particle, 600 is made to bind with three different oligonucleotides 602, 604 and 606. One of the oligonucleotides is bound to colloid particle 600 through a large complexing agent 608, e.g. a protein or a supramolecular structure, the purpose of which will be explained further below. In an array of three electrodes 610, each electrode is bound to one of double-stranded oligonucleotide 612, 614, and 616, each of which has a

different sticky end, complementary to a sequence in one of the oligonucleotides 602, 604 and 606. The oligonucleotides-dressed colloid particle is contacted with the electrode array and by providing conditions for hybridization, a structure 620, consisting of electrodes 610 bound to colloid particle 600 through three bridges, each of which consists of one of fibers 612, 614 and 616 and one of oligonucleotides 602, 604 and 606, respectively, is formed. Fibers 612, 614 and 616 may be made to covalently bind to oligonucleotides 602, 604 and 606, respectively, to form integral fibers 622, 624 and 626.

The colloid particle is subjected to passivation treatment, e.g. by means of exposure to alkane thiol, octadecyl thiol, dodecyl thiol, etc., to form an insulating layer 630 to isolate the colloid particle from the surrounding medium and avoid metal deposition on the colloid particle in a subsequent step. Fibers 622, 624 and 626 are then formed into wires, e.g. in the manner described above, whereby a SET is formed.

In the SET, wire 622 serves as a gate and for its proper function, a high resistance between it and the colloid particle 600 is preferred, which purpose is achieved by complexing agent 608. Under normal circumstances, colloid particle 600 resists a current flow, but when the potential at the gate is changed, the electrostatic field formed reduces the activation energy required to charge or discharge the colloid particle whereby current can flow between wires 624 and 626.

As will be appreciated, the manner of construction of the SET described herein is but an example. One alternative method is first formation of a colloid particle with fibers attached thereto and then causing the fibers to bind to the electrodes, e.g. through oligonucleotide linkers. As will be clear to the artisan, there are other possible alternatives allowing to construct a SET in accordance with the invention.

Reference is now being made to Fig. 7, illustrating a molecular switch which is based on a reversible photo transformation. A molecular fragment, such as bis thiophene derivatives of hexafluorocyclopentene or

maleimide, (Figs. 7A and 7B, respectively), are bound to the networks. Polymer groups (P1 and P2 (which may be the same or different)) that may contain recognition groups which may be sequence selective or non sequence selective, are attached to both thiophene moieties via covalent or non covalent interaction to form a disrupted conjugated polymer. The polymers P1 and P2 are each connected to a different fiber (P1 and P2, respectively – not shown). Upon exposure to light with an appropriate wavelength ( $\lambda$ ), photocyclization of the thiophenes with a double bond of the hexafluorocyclopentene or maleimide occurs, thus forming a conjugated polymeric wire electrically linking P1 and P2. Photo excitation of the cyclized switch results in the retrocyclization process and redisrupts the conjugation along the polymer.

The switching light signal may be provided from an external light source, or may be provided internally by any internal light source.

Fig. 8 is a schematic illustration of a FET in accordance with the invention. The FET 800 comprises source electrode 802, a drain electrode 804, situated at two ends of a semiconductor matrix 806. Gate wire 808, connected to gate electrode 810, is situated in a recess 812, within semiconductor matrix 806. The gate length in a semiconductor FET determines, to a large extent, the maximal frequency at which the FET can operate. Shorter gates minimize the electron flight time under the gate and hence facilitate higher operation frequencies. Since FET dimensions, apart from the gate, are not critical, the gate parameters set the bottle neck for mass production of faster FETS. The FET in accordance with the invention provides a solution to this problem. The FET, apart from the gate, may be fabricated by conventional lithography and semiconductor processes and techniques, and then the gate may be formed by stretching a nucleic acid fiber between electrodes and then the fiber may be metalized as outlined above. In this manner sub-0.1 micrometer gates can easily be formed allowing inexpensive mass production of faster FETS.

The invention will now be illustrated further in the following Examples.

## EXAMPLES

### Example 1: Preparation of Linkers

#### (a) Disulfide based linkers:

Controlled pore glass (CPG) derivatized with a disulfide group is used for the synthesis (starting from its 3' side) of an oligonucleotide having a free 5' site. The oligonucleotide is prepared using a conventional DNA synthesizer (see scheme in Fig. 9).

#### (b) Thiol-based linkers:

Linkers are being prepared according to (a) above and the disulfide bond is cleaved to obtain a free thiol.

#### (c) Biotin-streptavidin complex based linkers:

Biotin moiety is attached to an oligonucleotide having a specific sequence, as known *per se*. The biotin-oligonucleotide is coupled via a streptavidin molecule to another molecule containing a biotin moiety at one side (see also Fig. 1B) and a thiol or disulfide group on the other side.

#### (d) Repressor based linkers:

A nucleic acid binding protein, such as the *lac* repressor, is covalently attached to a thiol group. A DNA sequence is synthesized having sticky ends and containing the target sequence to which the repressor binds. The DNA sequence is coupled to the repressor through the target sequence.

#### (e) Thiophosphate based linkers:

The construction (starting from its 3' side) of an oligonucleotide sequence is carried out using a conventional DNA synthesizer wherein thiophosphates containing nucleotides are used instead of phosphate containing nucleotides.

#### (f) Artificial site specific based linkers:

A synthetic site-specific moiety such as, for example Rh(Phen)<sub>2</sub>Phi, known to bind 5'-pyr-pyp-pur-3' sequence<sup>(26)</sup> (pyr = pyrimidine, pur = purine), is covalently coupled to a thiol group.

Example 2: Attachment of the linker to an electrode

(a) Micropipette Wetting:

Electrodes are exposed to solutions of the appropriate linkers, for example, by employing pipettes or micropipettes or by any liquid dispensers. Such liquid dispensers may be fixed onto a manipulator that may be computer controlled. Different types of linkers can be deposited on each electrode. Additionally, different types of linkers can be deposited simultaneously or sequentially on different electrodes.

(b) Jet printing:

Ink-jet like printing techniques are used for the selective exposure of different electrodes to different linkers. By utilizing such a technique, it is possible to attain high precision, resolution and to increase rates of production, facilitating large scale production.

(c) Ab-initio electrode-linker synthesis:

(c1) Using selective masking techniques:

The well developed technology used for synthesizing DNA sequences may be harnessed for the *ab-initio* preparation of a complex electrode-linker array. For example: a substrate composed of a set of electrodes on an inert substrate is partially coated with an inert coating yielding two types of electrodes: coated electrodes (A) and uncoated electrodes (B). The substrate is exposed to a solution of a thiol linked to a nucleic acid sequence serving as a starting point for DNA synthesis. Due to the inert coating, only the uncoated B electrodes react with the thiol. Using standard DNA synthesizing techniques, a pre-defined sequence is produced on the B electrodes. The substrate is then rinsed and the masked electrodes are uncovered followed by the selective coating of B electrode. This procedure allows the production of two types of electrodes differing one from the other by the type of linkers bound thereto. The same technique with somewhat more complex steps (several steps of masking and unmasking) allows the fabrication of various substrates having many different electrodes with different linkers bound thereto.

(c2) Using photodeprotection techniques:

This approach involves the utilization of photolabile groups for the protection of the start point of DNA synthesis. Inactivated start point groups are unable to react with nucleotides. Using selective irradiation by means of a mask and/or a light conductor and/or any other addressable light source, the activation of different selected electrodes is achieved by the photoremoval of protecting groups from selected electrodes.

(c3) Using blockers:

Using the masking technique ((c1) above) a set of electrodes is prepared for oligonucleotide synthesis. Once a DNA sequence is completed on one set of electrodes, a terminating group is attached to the oligonucleotide ensuring their inertness. Other sequences can be further synthesized on different electrodes that are prepared according to the previous step but become active according in this step. It should be noted that the set of linkers constructed in the previous step is not affected due to the attached blockers to their end points.

(c4) Electrode printing:

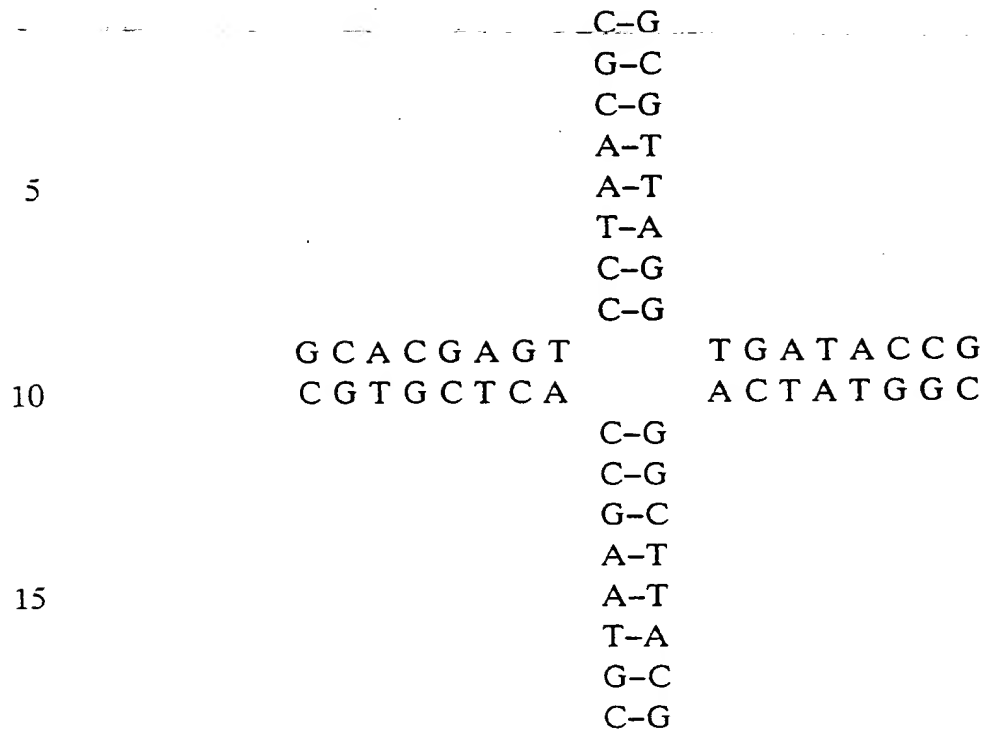
Linkers are attached to conducting beads such as gold colloids. The colloids are then dispersed in a controllable manner to form conducting metal pads with linkers attached thereto. Dispensing may be achieved by the different techniques outlined above or by any conventional technique.

The above techniques may be used alone or in any combination with other techniques.

**Example 3: Construction of networks - Production of Junctions**

(a) Production of a branched sequence:

A stable four-arm branched DNA junction is constructed using for example the following sequences:



20

Careful planning of the sequences allows the fabrication of complex junctions according to a desired design. This branch sequence may be attached to double-stranded fibers using methods known *per se*.

(b) Creating a branch by enzymatic reactions:

25

The protein *recA* from *E. coli* bacteria catalyzes the recombination and construction of a base-paired hybrid to joining two DNA molecules. It can join, in a specific way, a single-stranded DNA with a double-stranded DNA provided that homology exists between the single-stranded and the double-stranded DNA. DNA-binding proteins can extend single-stranded DNA and facilitate DNA annealing by random collisions. It is also possible to achieve base-pair specific contacts between two separate duplexes to form a four-stranded structure that is aligned through chemical moieties exposed in the grooves of the two helices. Similarly, the *recA* protein can induce specific contacts between a single-stranded and a double-stranded DNA, through recognition of the complementary sequence from the "outside" without the need to open the double strands and expose

30

35

to the single-strand sequence. There is also the possibility to recombine three-stranded and four-stranded DNA helices. Four DNA strands can also undergo "switch pairing" at a joining point to form a crossed-strand junction (called *Holliday* structure). There is then the possibility to create the so called heteroduplexes which are regions on recombinant DNA molecules where the two strands are not exactly complementary. The branching joint, however, can migrate to its equilibrium point of complementary base pairing. Fully recombinant duplexes are formed by allowing steric rearrangements. The utilization of *RecA* protein for making the Holliday structures *in vitro* is well established (see, for example B. Alberts, *et al.*, *Molecular Biology of the Cell*, 3rd Edition, Garland Publishing Inc., New York, 1994). Another enzyme, *RecBC* from *E. coli* bacteria has both unwinding and nuclease activities and can therefore catalyze the exposure of single-stranded DNA with a free end, allowing the *RecA* protein to start the pairing reaction. Important for the step-by-step build-up of the network, is the fact that *RecBC* initiates unwinding only on DNA containing a free duplex end. It then navigates along the DNA, from the free end, unwinding and rewinding DNA as it goes. Because it unwinds the DNA faster than rewinding it, "bubbles" or loops of single-stranded regions are created in the duplex DNA. *RecA* protein can then bind to a cut, made by the *RecBC* in a specific sequence, 5'-GCTGGTGG-3', in one of the single strands and initiates DNA strand exchange with another DNA. The specific sequence of the cut can be pre-designed by artificial recombinant DNA synthesis (see B. Alberts *et al.*, *supra* and L.P. Adams *et al.*, *The Biochemistry of the Nucleic Acids*, 11th Edition, Chapman & Hale, 1992).

(c) Utilizing nucleic acid-binding proteins:

Two or more specific DNA binding proteins are allowed to interact with two or more DNA strands. Coupling of such binding proteins enables the formation of a junction between the different DNA strands.

**Example 4: Connection of network to substrate**

Anchoring the network to the substrate may be realized using various DNA binding proteins. For example, repressors from bacteria (e.g. *lac*-repressor or  $\lambda$ -repressor) which can bind to both the substrate (such as a plastic substrate) and to the DNA thus joining the two. Such binding stabilizes the network without necessarily taking part in the electrical functionality.

**Example 5: Preparation of an integrated circuit**

The integrated circuit (IC) is composed of a substrate such as silicon, derivatized silicon, silica, derivatized silica, organic polymer or any other substance capable of acting as a support for the fabrication or mechanical fixation or stabilization of the network. The substrate may serve an electrical function.

A typical example for IC preparation is described in the following:

**Example 6: Passivation of a glass substrate**

A glass substrate is immersed in fuming nitric acid (100%  $\text{HNO}_3$ ) for 10 min, rinsed in deionized (DI) water, then immersed in 1 N NaOH solution for an additional 10 min and rinsed in DI water. The cleaned glass is dried thoroughly, then immersed for c.a. 12 hrs in a solution of an alkyl trichlorosilane (octyl trichlorosilane, *t*-butyl trichlorosilane etc.) in tetrachloroethane (1:5 v/v). The glass plate is then rinsed carefully several times with tetrachloroethane and isopropanol, then dried thoroughly.

**Example 7: Electrode fabrication**

Electrodes are fabricated according to one of the following routes:

- (i) Standard photo, electron, or x-ray lithography on the substrate and subsequent deposition of conductive substance (e.g. metal). Alternatively, the conductive substance may be deposited first and patterned next.
- (ii) Electrode assembly onto the surface: Patterning of the glass surface using

polyelectrolytes such as polyethyleimine, polyalcoholes, polyacids, polypyridines etc. or other ligating agents such as a thiol monolayer (fabricated from organic compounds containing thiol and silane moieties at opposite sites on the molecular skeleton) followed by the fixation of electrically conducting components such as Gold colloids enabling the assembly of conducting electrodes onto the substrate.

**Example 8: Electrical functionalization of the nucleic acid based networks – metal based conductive wires**

(i) The relevant part of the network is exposed to a solution containing the appropriate metal ion, thus, ion exchange occurs at the phosphate groups of the DNA skeleton exposed to the solution. Intercalation of ions inside the DNA may also take place under certain conditions

(ii) The ion exchanged DNA complex is then reduced by a reducing agent such as hydroquinone.

Cycles (i) and (ii) can be repeated in a sequential order until a conducting wire is achieved. Alternatively, the formation of conducting metal wire includes the following steps as stand-alone processes or in conjunction with steps (i) and (ii) or combined with one or more of the following techniques.

(iii) The relevant part of the ion-exchanged network is exposed to a metastable mixture of the reducing agent and the metal ion. Reduction takes place only at the surface of the metal clusters formed by steps (i) and (ii) thus, the gap between the metal clusters is bridged by the metal deposition process.

(iv) The ion exchanged DNA or the partially treated DNA network is exposed to electrochemical process, transforming the ions loaded on the DNA polyelectrolyte into a metallic conductor. In addition, electrochemical processes along the DNA molecule promote the vectorial growth of the metal wire along it.

(v) Photochemical deposition of the metal from its corresponding ions for the formation of the metallic wire.

(vi) Clusters or colloids are adsorbed onto the DNA network using sequence selective components, for example, specific sequences which are capable of binding to specific sites on the DNA non-sequence-specific binding agents, e.g. polyelectrolytes undergoing electrostatic interactions with the DNA. These Clusters and/or colloids serve as catalysts for processes (iii)-(v) above.

(vii) Defects in granular wires fabricated by one or more of the above techniques may be annealed using diverse methods such as thermal annealing processes, electrodeposition, etc.

An example of the fabrication of a silver-functionalized network is as follows:

(i) A DNA network fixed on a substrate is exposed to a basic solution of silver ions ( $\text{pH}=10.5$ ,  $\text{NH}_4\text{OH}$ ,  $0.1 \text{ M AgNO}_3$ ). After the DNA polyelectrolyte is exchanged by the silver ions, the substrate is rinsed carefully with deionized water (DI) and dried.

(ii) The silver loaded DNA network fixed on a substrate is exposed to a basic solution of hydroquinone ( $0.05 \text{ M}$ ),  $\text{pH} = 5$  as a reducing agent. Steps (i) and (ii) are repeated sequentially until an electrically conducting wire is formed.

Complementary processes:

(a) step (iii) is performed after one or more (i)+(ii) cycles.

(iii) The DNA network loaded with silver metal clusters (after cycles (i) and (ii) have been performed) and after final rinsing with DI water is exposed to an acidic solution of hydroquinone (citrate buffer,  $\text{pH}=3.5$ ,  $0.05 \text{ M}$  hydroquinone) and  $\text{AgNO}_3$  ( $0.1 \text{ M}$ ). Cycle (iii) is terminated when the wire width attains the desired dimension. The process can be made light sensitive and thus can also be controlled by the illumination conditions.

(b) Electrochemical deposition for improved process:

(iv) In order to improve the aspect ratio of the metallic conductor, an electrochemical process is performed. For that purpose, pre-treatment with an alkane thiol is performed prior to the (i)+(ii) processes. This ensures the inertness of the metal electrodes against electrochemical metal deposition.

After one or more of the (i)+(ii) cycles, the electrodes connected through the DNA covered metal wire are connected to a current and bias controlled electrical source and the relevant part of the DNA network is exposed to a solution of the metal ion (different concentrations according to a specific protocol). The gaps between the conducting domains are via electrochemical metallic deposition.

(c) Photochemical deposition for an improved process:

(v) In order to improve the aspect ratio of the metallic conductor, a photochemical process is performed in a similar manner to the electrochemical process outlined above but using photochemical reaction as driving processes. For example, metalization of a DNA network may be obtained using an electron donor (triethanolamine, oxalic acid, DTT etc.), a photosensitizer (Ru-polypyridine complexes, xanthene dyes semiconductor particles such as  $\text{TiO}_2$ , CdS etc.), an electron relay such as different bipyridinium salts and the relevant metal ion or metal complex. The photosensitizer transduces the absorbed light energy into a thermodynamic potential through electron transfer processes involving the electron donor and electron acceptor in any of the possible sequences. The reduced electron acceptor acts as an electron relay and charges the metal clusters/colloids with electrons. The charged clusters/colloids act as catalysts for the reduction of the metal ions thus inducing the growth of the metal conductor.

(d) Gold clusters and/or colloids as nucleation centers:

(vi) Instead of performing the first (i)+(ii) cycles, the relevant part of the DNA network is exposed to a solution of gold colloids pre-coated (partially) with cationic thiols (such as pyridinium alkane thiol). The Gold colloids are being adsorbed to the DNA skeleton by ion pairing and the growth of the wire is attained using one or more of the above techniques.

(e) Curing processes:

(vii) Defects in a granular wire obtained by one or a combination of the above techniques are annealed using diverse processes such as thermal annealing processes (hydrogen atmosphere (10%  $\text{H}_2$  in  $\text{N}_2$ ), 300C over several hours).

**Example-9: Connecting two electrodes by a conductive wire formed on a DNA template, and properties of the wire**

(a) Wire preparation:

- 5                Figure 3A outlines the DNA templated assembly of the metal wire. A glass coverslip was first passivated against spurious DNA binding. Subsequently, two parallel gold electrodes were deposited on the coverslip using standard microelectronic techniques. One gold electrode was then wetted with a micron size droplet of an aqueous solution containing a 12-
- 10    base, specific sequence oligonucleotide, derivatized with a disulfide group attached to its 3' side. Similarly, the second electrode was marked with a different oligonucleotide sequence. After rinsing, the sample was covered by a solution of  $\sim 16 \mu\text{m}$  long  $\lambda$ -DNA, having two 12-base sticky ends that were complementary to the oligonucleotides attached to the gold electrodes.
- 15    A flow normal to the electrodes was induced to stretch the DNA, allowing its hybridization with the two distance surface-bound oligonucleotides. Stretching the DNA between two electrodes could also be carried out in reverse order, where hybridization and ligation of the disulfide derivatized oligonucleotides to the long DNA molecule was performed prior to its
- 20    application to the sample. Both methods work equally well. Fig. 10 depicts a fluorescently-labeled  $\lambda$ -DNA bridging two gold electrodes. By observing the curving of the DNA molecule under perpendicular flow it was demonstrated that it was attached solely to the electrodes. Sample preparation was completed by removal of the solutions.
- 25                Two-terminal measurements performed on these samples prove that the stretched DNA molecule was practically an insulator with a resistance higher than  $10^{13} \Omega$ . The insulating nature of the DNA was in accordance with previous spectroscopic electron-transfer rate measurements<sup>(18)</sup>. To instill electrical functionality, silver metal was vectorially
- 30    deposited along the DNA molecule. The three-step chemical deposition process was based on selective localization of silver ions along the DNA through  $\text{Ag}^+/\text{Na}^+$  ion-exchange<sup>(19)</sup> and formation of complexes between the silver and the DNA bases<sup>(19-22)</sup>. The  $\text{Ag}^+/\text{Na}^+$  ion-exchange was monitored

by following the quenching of the fluorescence signal of the labeled DNA. The process was terminated when full quenching was achieved. After rinsing, the silver ion-exchanged - DNA complex was reduced using a basic hydroquinone solution. This step resulted in the formation of nanometer  
5 size metallic silver aggregates bound to the DNA skeleton. These silver aggregates serve as spatially localized nucleation sites for subsequent growth of the wire. The ion-exchange process was highly selective and restricted to the DNA template only. The silver aggregates, bound to the DNA, were further "*developed*", much as in the standard photographic procedure, using  
10 an acidic mixture of hydroquinone and silver ions under low light conditions<sup>(24,25,32-37)</sup>. Acidic solutions of hydroquinone and silver ions are metastable but spontaneous metal deposition is normally very slow. The presence of metal catalysts (such as the silver nucleation sites on the DNA), significantly accelerates the process. Under these experimental conditions,  
15 metal deposition therefore occurs only along the DNA skeleton, leaving the passivated glass practically clean of silver. The process was terminated when the trace of the metal wire was clearly observable under a differential interference contrast (DIC) microscope. The metal wire followed precisely the previous fluorescent image of the DNA skeleton. The structure, size and  
20 conductive properties of the metal wire were reproducible and dictated by the "*developing*" conditions.

#### Results:

Atomic force microscope (AFM) images of a section of a 100 nm wide, 12  $\mu\text{m}$  long wire are presented in Fig. 11. As clearly seen, the wire  
25 comprises of 30-50 nm-diameter grains deposited along the DNA skeleton.

To study the electronic transport properties of these wires, two terminal I-V curves have been measured at room temperature using an HP parameter analyzer with internal resistance of  $10^{13}\Omega$  and current resolution of 10 fA. Figure 12 A shows the I-V curves of the silver wire presented  
30 in Fig. 11. The curves are highly non linear and asymmetric with respect to zero bias. The shapes of the curves depend on the voltage scan direction indicated by arrows in Fig. 12A. Approaching zero voltage from large

positive or negative bias, the current vanishes almost linearly with voltage.

A zero-current plateau then develops with differential resistance larger than  $10^{13}\Omega$ . At higher bias, the wire turns conductive again with a differential resistance somewhat lower than in the original bias polarity. Repeated measurements in the same scan direction (solid curves in Fig. 12A) yield reproducible I-V curves. The length of the zero bias plateau in different wires can vary from a fraction of a volt to roughly 10 volts. The solid line in Fig. 12B depicts, for example, the I-V curve of a different wire in which the silver growth on the DNA was more extensive. As a result, the plateau was reduced to 0.5 volts. By driving higher currents through the wire, the plateau could usually be eliminated to give an ohmic behavior (dashed line in Fig. 12B) with resistance varying between 1 and 30 M $\Omega$ , depending on the silver deposition process.

#### 15    **Example 10: Organic conjugated-polymer based conducting wires**

(i) The relevant part of the network is exposed to a solution containing a cationic segment capable of forming a conjugated-polymer by a chemical transformation or a cationic non conjugated-polymer capable of undergoing conjugation by a chemical transformation or a cationic conjugated-polymer. Thus, ion exchange process occurs at the phosphate groups of the DNA skeleton exposed to the solution.

(ii) The ion exchanged DNA complex is treated according to the nature of the organic species that is bound to the polyanionic skeleton. Electrical functionalization is achieved either by the former process or by a sequential doping process. Doping may be achieved via conventional redox processes, by protonation - deprotonation processes, by electrochemical means or by photochemical means. Additionally, sequence selective processes between the DNA skeleton and the building blocks of the above organic conjugated-polymer based conducting wires can be utilized for the production of wires with a well defined structure, electrical affinity gradients and p/n junctions.

I. The fabrication of a PPV (poly-phenylene vinylene) conducting wire is as follows:

(i) A DNA network fixed on a substrate is exposed to a solution of a pre-PPV water soluble polymer. After the DNA polyelectrolyte is ex-  
5 changed by the pre-PPV polymer, the substrate is rinsed carefully and dried.

(ii) The pre-PPV polymer loaded DNA network fixed on the substrate is reacted in a vacuum oven (e.g  $1\text{e-}6$  bar, 300 C, 6hr.).

(iii) The resulting luminescent PPV polymer is doped using conventional methods until displaying conductivity.

10 II. An alternative route for the fabrication of a PPV conductive wire is as follows:

(i) A DNA network fixed on a substrate is exposed to a solution of a bis-(tetrahydrothiophenium)-p-xylylene dichloride. After the DNA polyelectrolyte is exchanged by the bis-(tetrahydrothiophenium)-p-xylylene  
15 dichloride, the substrate is rinsed carefully and dried.

(ii) The bis-(tetrahydrothiophenium)-p-xylylene dichloride loaded DNA network fixed on a substrate is polymerized in a basic solution to form a pre-PPV polymer attached to the DNA backbone.

(iii) The pre-PPV polymer loaded DNA network fixed on a substrate  
20 is reacted in a vacuum oven ( $1\text{e-}6$  bar, 300 C, 6hr.).

(iv) The resulting luminescent PPV polymer is doped using conventional methods until displaying desired conductivity.

III. The fabrication of a PANI (polyaniline) conducting wire is carried out as follows:

25 (i) A DNA network fixed on a substrate is exposed to a solution of an acid soluble PANI polymer. After the DNA polyelectrolyte is exchanged by PANI polymer, the substrate is rinsed carefully and dried.

(ii) The resulting PANI polymer is doped using conventional methods until displaying desired conductivity.

30 IV An alternative route to the fabrication of a PANI conducting wire is as follows:

(i) A DNA network fixed on a substrate is exposed to a solution of anilinium ions. After the DNA polyelectrolyte is exchanged by the anilinium ion, the substrate is rinsed carefully and dried.

5 (ii) The anilinium ions loaded on the DNA skeleton are oxidized using a solution of an oxidizing agent such as peroxodisulphate ions, yielding a polyaniline polymer. The resulting PANI polymer is doped using conventional methods until displaying desired conductivity.

V. An alternative route to the fabrication of a PANI conducting wire is as follows:

10 (i) A DNA network fixed on a substrate is exposed to a solution of a short oligomer of PANI (>1 repeat unit). After the DNA polyelectrolyte is exchanged by the PANI oligomer, the substrate is rinsed carefully and dried.

15 (ii) The PANI oligomer ions loaded on the DNA skeleton are oxidized using a solution of an oxidizing agent such as peroxodisulphate ions, yielding a polyaniline polymer. The resulting PANI polymer is doped using conventional methods until displaying desired conductivity.

#### **Example 11: Fabrication of insulators**

20 Insulators may be constructed on electrically functionalized parts of the network such as wires and connections between wires and components.

##### **A. Insulation of metallic components (including metallic wires):**

25 (1) Metallic components may be electrically insulated from their environment using surface active agents composed of an insulating body such as an alkyl group attached to a surface binding group such as a thiol or a disulfide group. The surface binding group binds to the surface of the metal thus, forming a dense *"two dimensional"* layer of electrically  
30 insulating molecules at the surface of the metal. This layer presents an electrical barrier, namely, an insulator.

(2) The selective oxidation and other derivations that form non conducting layers and interfaces form electrically insulating barriers too.

**B. Insulation of non metallic components (including conjugated-polymer based wires):**

Using complementary interaction based molecular recognition processes, insulating layers that can self assemble onto the surface of non metallic components are constructed. For example, positively charged polymers such as PANI can be coated and insulated from their environment using a polyanion polymer having long alkyl chain side groups.

**Example 12: Preparation of a single electron transistor (SET)**

In the following the main steps in the fabrication of a SET are outlined:

1. Substrate passivation is carried out according to Example 6.
2. Gold electrode definition by standard microelectronic techniques is carried out, e.g. as specified in Example 7. For the simple case of a single SET only three electrodes are needed.
3. Linker synthesis, e.g. carried out according to Example 1(a). The Linkers are typically made of short (for example 12 bases) oligonucleotides derivatized with a disulfide group at their 3' side. For the case of a single SET, linkers with three different sequences are preferable.
4. Linkers are made to interact with the electrodes through their disulfide group. Each electrode is marked by a different linker according to Example 2.
5. Three different disulfide derivatized linkers are attached to the same nanometer scale gold colloid at their disulfide side and one of the linkers is attached via a large bridging agent which creates a large barrier between the colloid and this linker.
6. Three double-stranded DNA molecules, a few micron long, containing sticky ends complementary to those of the linkers at the gold colloid are hybridized and ligated with the linkers attached to the colloid to form a

three terminal junction with the colloid at the branching point. The three DNA molecules also contain sticky ends complementary to the three linkers attached to the electrodes, at their sides opposite to the gold colloid.

7. The network is completed by letting the double-stranded DNA molecules attached to the gold colloid to hybridize with the three linkers on the electrodes, followed by ligation of the nicks.

8. Electrical functionality is achieved by the following steps.

a) The gold colloid is coated by a thiol group layer to be used as an insulating barrier, and passivation layer against metal deposition.

10 b) The DNA fibers are coated with metal to form conducting wires, for example as specified in Example 8. The wires are electrically weakly coupled to the gold colloid through the thiol group insulating barrier with the wire coupled through the large bridging agent having even weaker coupling to the colloid.

15 c) Fig. 6 outlines the final device composed of a Gold colloid wired to three electrodes through the insulating barriers.

The current between two electrodes can now be modulated by a small voltage applied to the third electrode (the one very weakly coupled to the colloid). The circuit hence function as a SET. The SET can be fabricated as a part of a more complex circuit where the electrodes are replaced by functionalized network components.

It should be appreciated that the sequence of steps from the preparation of the SET may be altered, thus for example, step 6 may precede step 4, etc.

25

#### **Example 13: PPV functionalized fiber as a light source**

The process described in Example 10 may be followed up to and including step I(ii). The resulting PPV component is highly luminescent and has a width considerably smaller than 100 nm. Fabricating the PPV component between electrodes of appropriate work functions then forms an electroluminescent device.

30

**CLAIMS:**

1. A microelectronic circuit comprising a network of at least one electric or electronic component, the network being connected to at least one, interface components, characterized in that  
5 the network has a geometry defined by at least one nucleic acid fiber; each of said interface components is connected to at least one fiber; and in that  
said at least one fiber has one or more substances or particles bound thereto or complexed therewith such that at least one electric or electronic component is formed with properties based on electric charge transport characteristics of said one or more substance or particles.  
10
2. A microelectronic circuit according to Claim 1, wherein said properties are based also on location and connectivity of said at least one electric or electronic component within the network.  
15
3. A microelectronic circuit according to Claim 1 or 2 wherein the network is connected to at least two interface components.
4. A microelectronic circuit according to any one of Claims 1-3, wherein the network has at least one conductor and at least one microelec-  
20 tronic component, characterized in that
  - (i) the conductor is a wire constructed on at least one first portion of the at least one nucleic acid fiber which is electrically functionalized by having one or more electrically conductive substances or particles bound thereto such that electric current can flow along said at least one first portion of the nucleic acid fiber;  
25
  - (ii) the microelectronic component being electrically connected to said at least one wire and being constructed either on at least one second portion of a nucleic acid fiber, other than said at least one first portion, by having one or more substances or particles bound thereto rendering said at least one second portion to have  
30 electronic functionality, or being constructed by a particle situated at a junction between two or more nucleic acid fibers rendering

~~said junction to assume an electronic functionality, said electronic functionality being based on electric charge transport characteristics of said one or more substances or particles;~~

(iii) each of said input/output interface components is connected to at least one functionalized nucleic acid fiber or at least one of said micro-electronic components in a manner to allow electric conductivity between said interface component and said functionalized fiber or said micro-electronic component, respectively.

5  
10 5. A microelectronic circuit according to any one of Claims 1-4, wherein the network comprises nucleic acid fibers which are assembled to form a network on the basis of their sequence specific hybridization with other fibers or specific binding to other components.

6. A microelectronic circuit according to any one of Claims 1-5, wherein the network comprises at least one fiber electrically or electronically functionalized by binding one or more of said substances or particles thereto on the basis of non-sequence selected molecular recognition properties of the fibers.

7. A microelectronic circuit according to any one of Claims 1-5, wherein the network comprises at least one fiber electrically or electronically functionalized by said substances or particles bound to the fiber at specific domains or sequences of the fiber.

8. A microelectronic circuit according to any one of Claims 1-7, wherein the network comprises at least one wire fabricated from conducting metal substances bound to the fiber or portion thereof.

25 9. A microelectronic circuit according to any one of Claims 1-7, wherein the network comprises at least one wire formed by non-metallic conducting substance bound to a fiber or portion thereof.

10. A microelectronic circuit according to any one of Claims 1-9, wherein the network comprises particles bound at junctions between fibers.

30 11. A circuit according to Claim 10, wherein said particle is a colloid particle.

12. A circuit according to Claim 11, wherein the junction comprising the colloid particle functions as a single electron transistor (SET).

13. A circuit according to any one of Claims 1-12, wherein the network comprises at least one fiber having at least a portion bound to  
5 semi-conducting substances.

14. A circuit according to Claim 13, wherein adjacent portions of a fiber are bound one to a p-type semi-conducting substance and the other to an n-type semi-conducting substance, whereby the two adjacent portions of the fiber constitute a p/n function.

10 15. A circuit according to any one of Claims 1-14, wherein the network comprises at least one nucleic acid-based junction formed by hybridization of complementary sequences in at least two fibers.

16. A circuit according to Claim 15, wherein at least one of the nucleic acid-based junctions is bound to either an n-type semi-conducting  
15 substance or a p-type semi-conducting substance with portions flanking the junction being deposited by the other of the n- or p-type semi-conducting substance, whereby the junction constitutes a bipolar transistor.

17. A component useful in a network as defined in any one of Claims 1-16.

20 18. A component according to Claim 17, selected from the group consisting of a switch, bipolar transistor, single-electron transistor, field effect transistor, diode, capacitor, resistor and a conductor.

19. A wire comprising a nucleic acid fiber having bound thereto an electrically conducting substances such that electric current can flow along  
25 the fiber.

20. A wire according to Claim 19, wherein the electrically conducting substance is metal, non-metal conductor or any combination thereof.

21. A field effect transistor comprising a wire according to Claim 19 or 20 serving as its gate.

22. A component according to Claim 17, comprising a fiber having poly-phenylene vinylene bound thereto, constituting a light producing component.

23. A method for fabricating a microelectronic network comprising

5 (a) providing an arrangement consisting of at least one interface components;

(b) immobilizing a linker on said interface component capable of binding to a nucleic acid sequence,

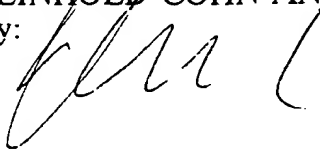
10 (c) contacting said arrangement with at least one nucleic acid fiber with a sequence capable of binding to the immobilized linker agent, and permitting binding of said sequence to said linker,

15 (d) functionalizing the at least one fiber by depositing thereon or complexing thereto at least one substance or particles imparting electric or electronic functionality to the fibers, said functionality of said electric or electronic component being based on electric charge transport characteristics of said at least one substance or particles.

24. A method according to Claim 23, wherein the network is formed by self-assembly as a result of chemical complementary and molecular recognition properties of at least one fiber to at least one other fiber or 20 between at least one fiber and at least one specific sequence- or domain-recognizing complexing agent.

25. A method according to Claim 23 or 24, comprising mixing fibers and components together and allowing them to self-assemble into a network in a specific manner.

25 26. A method according to any one of Claims 23-25, comprising forming junctions between fibers with particles situated in the functions, said particles serving as an electronic component in the network.

For the Applicants,  
REINHOLD COHN AND PARTNERS  
By: 

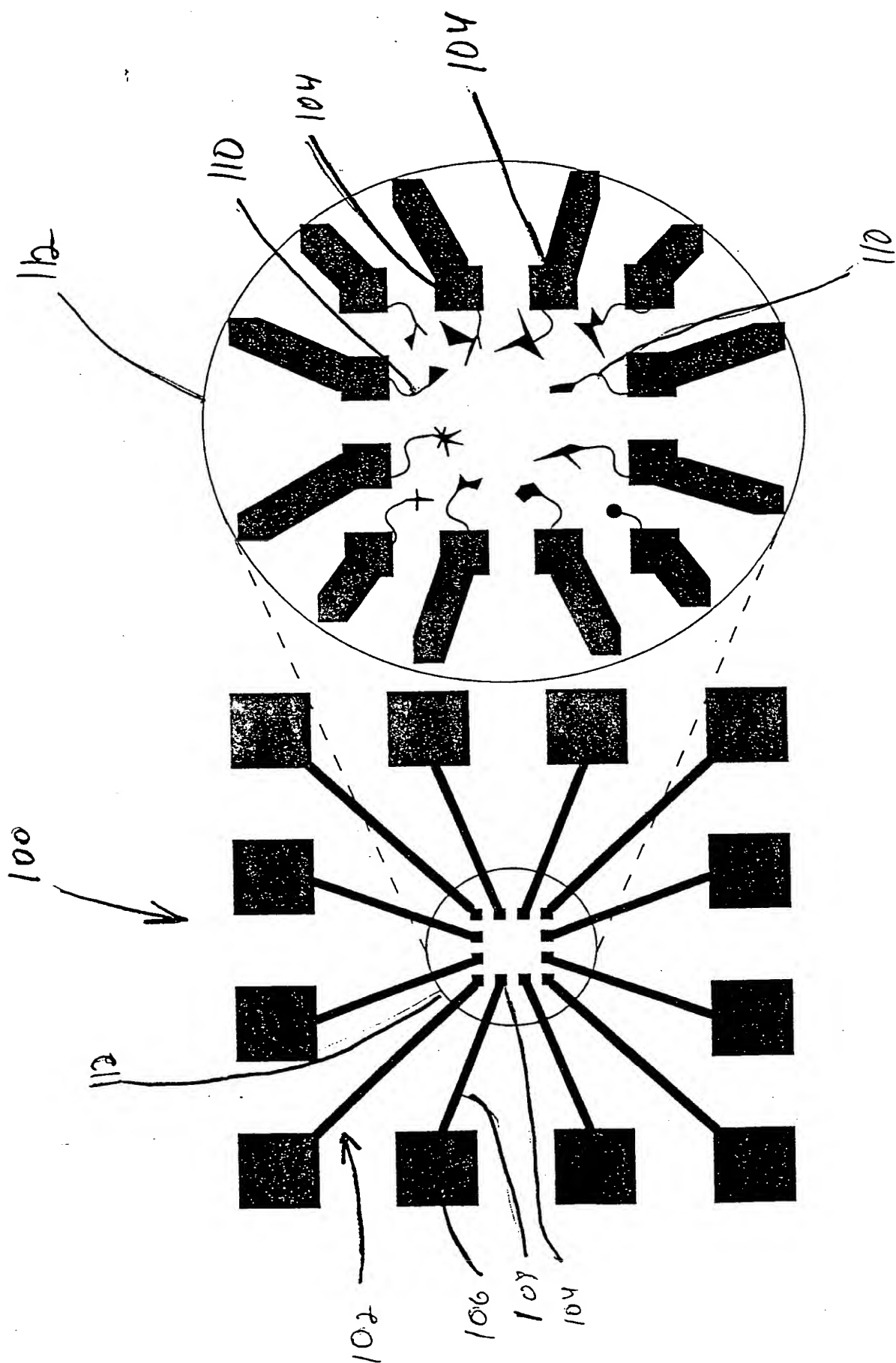


Fig. 1A

~~Scheme 1~~

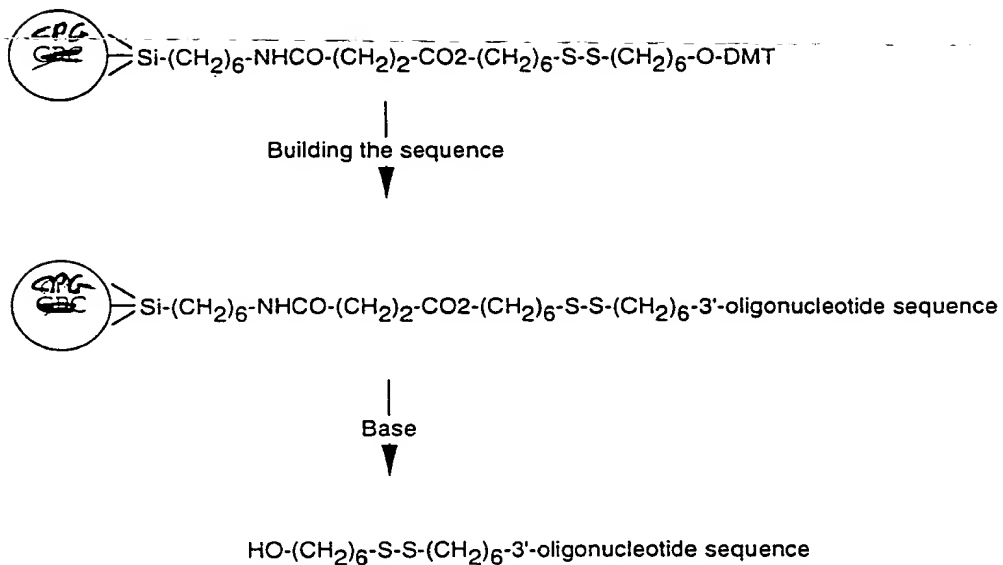


Fig. 9

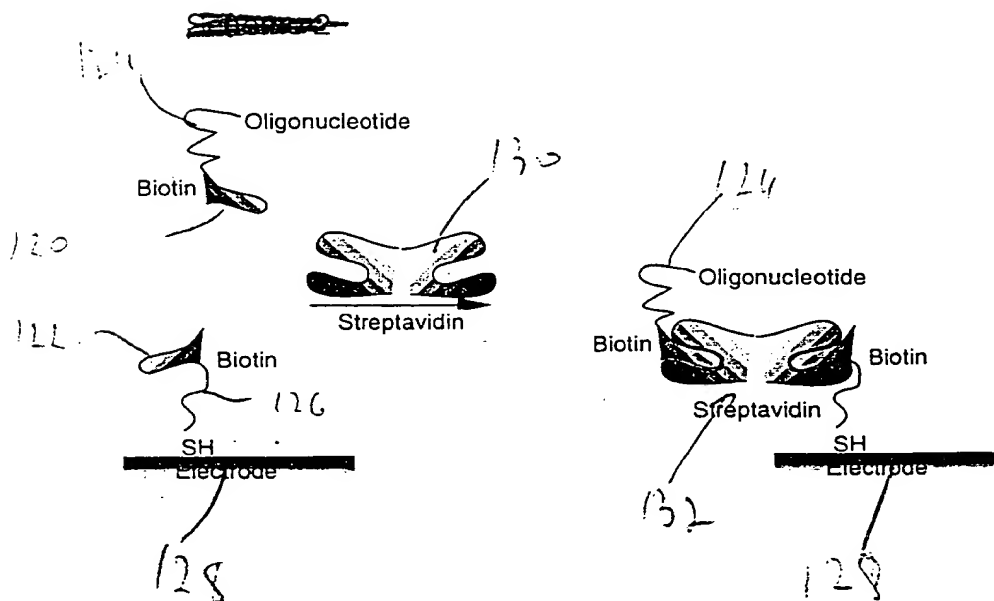


Fig. 1.B

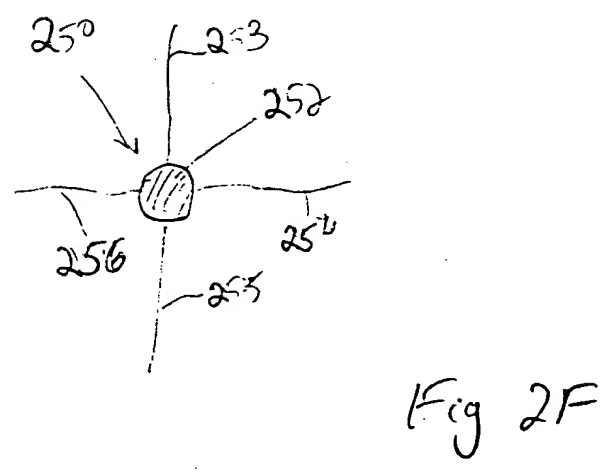
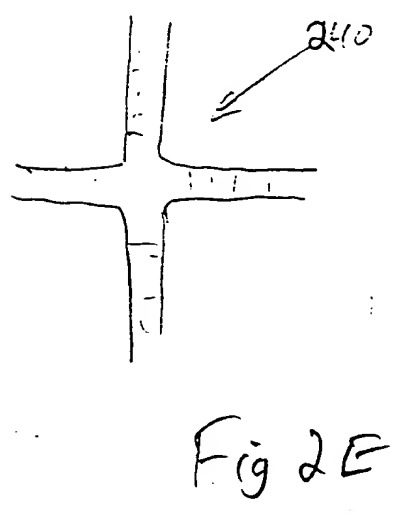
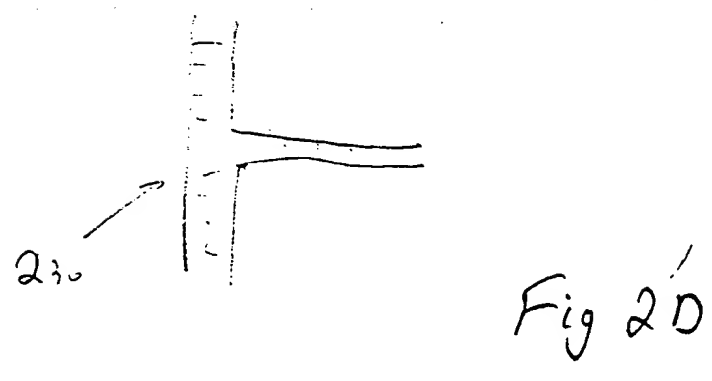
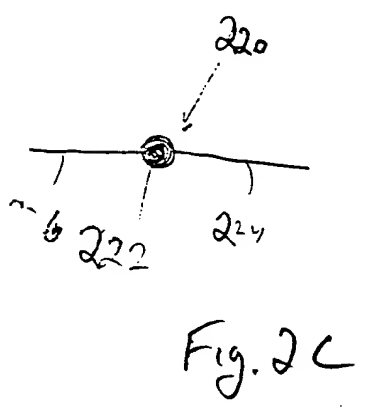
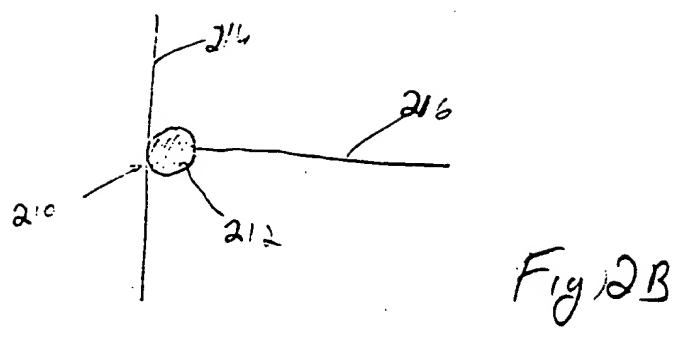
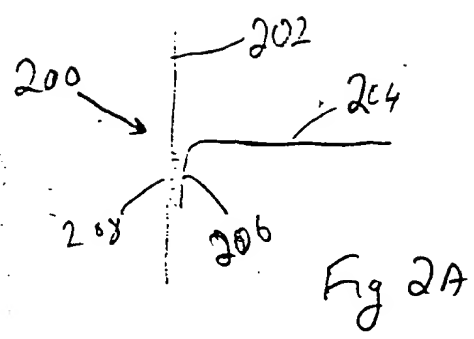


Fig. 2

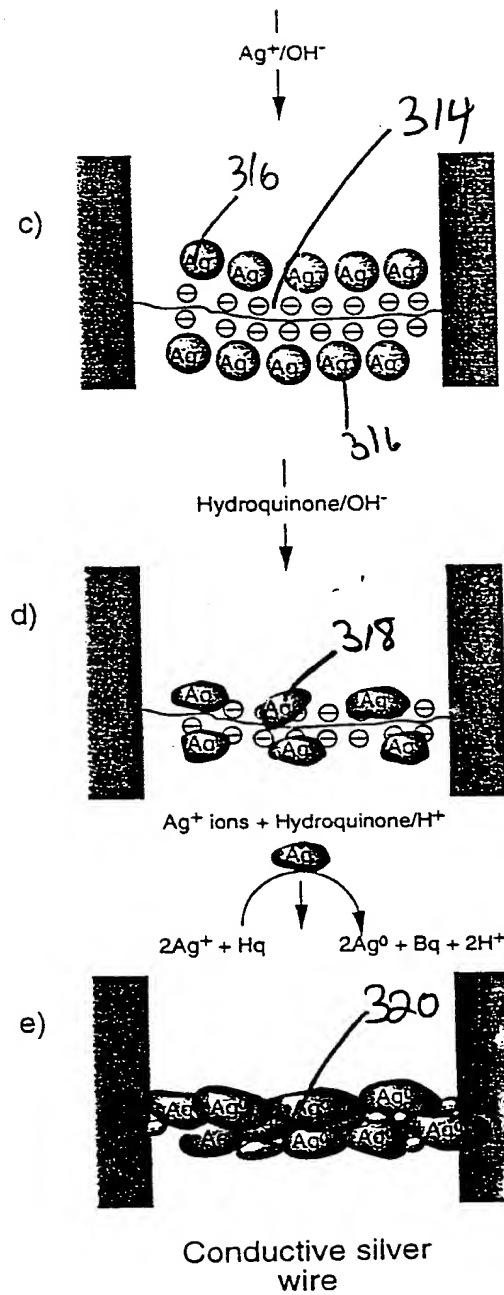
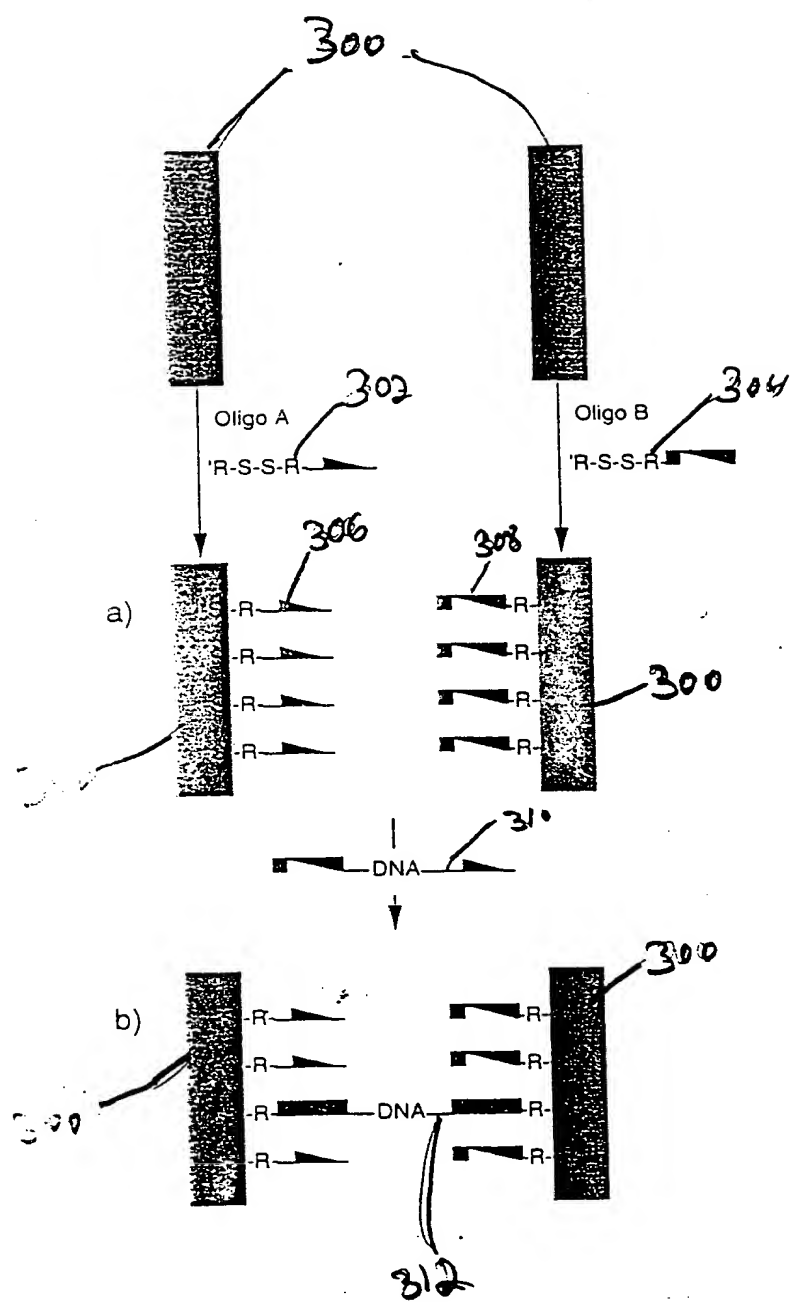


Fig. 3A

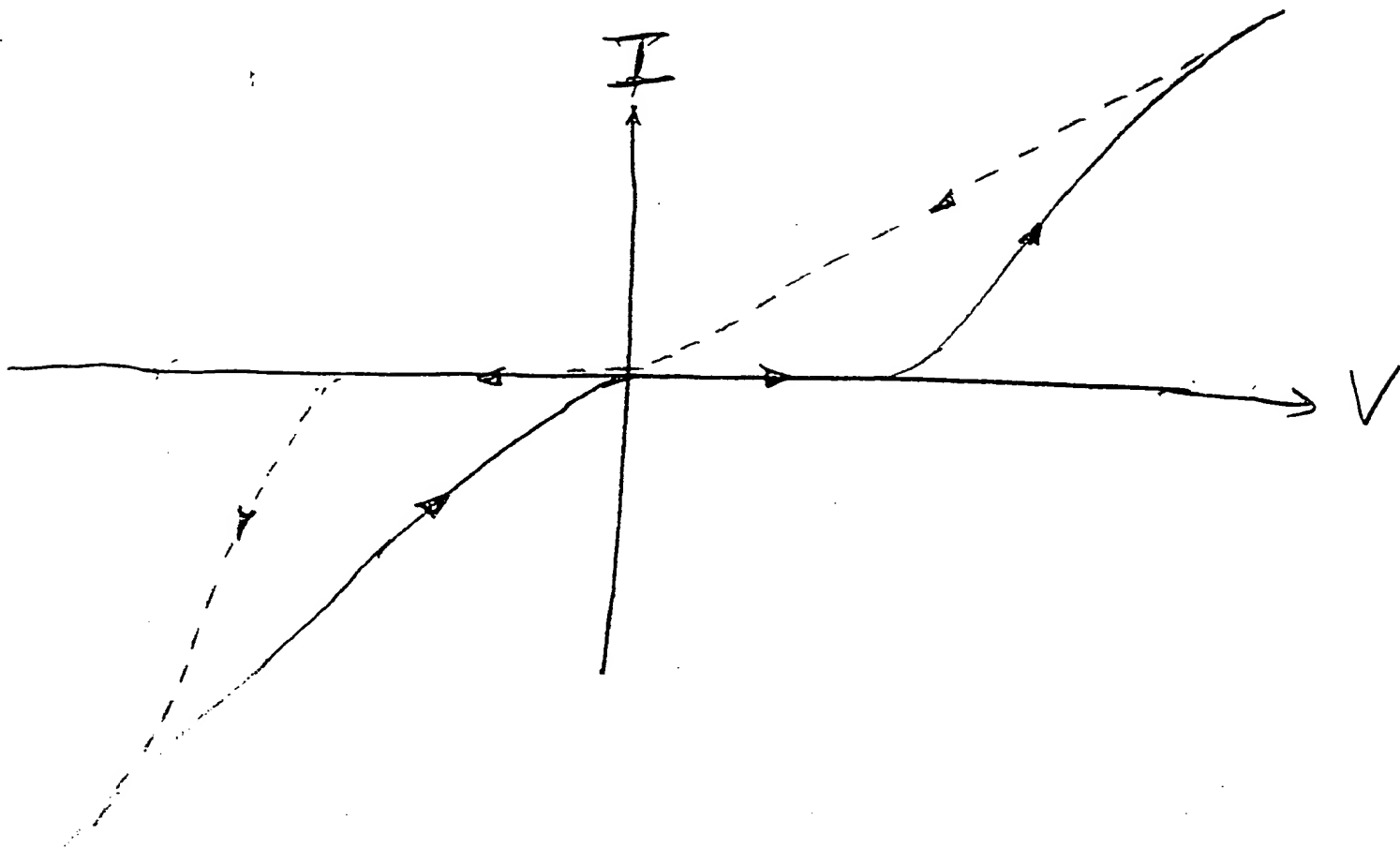


Fig 3B

7

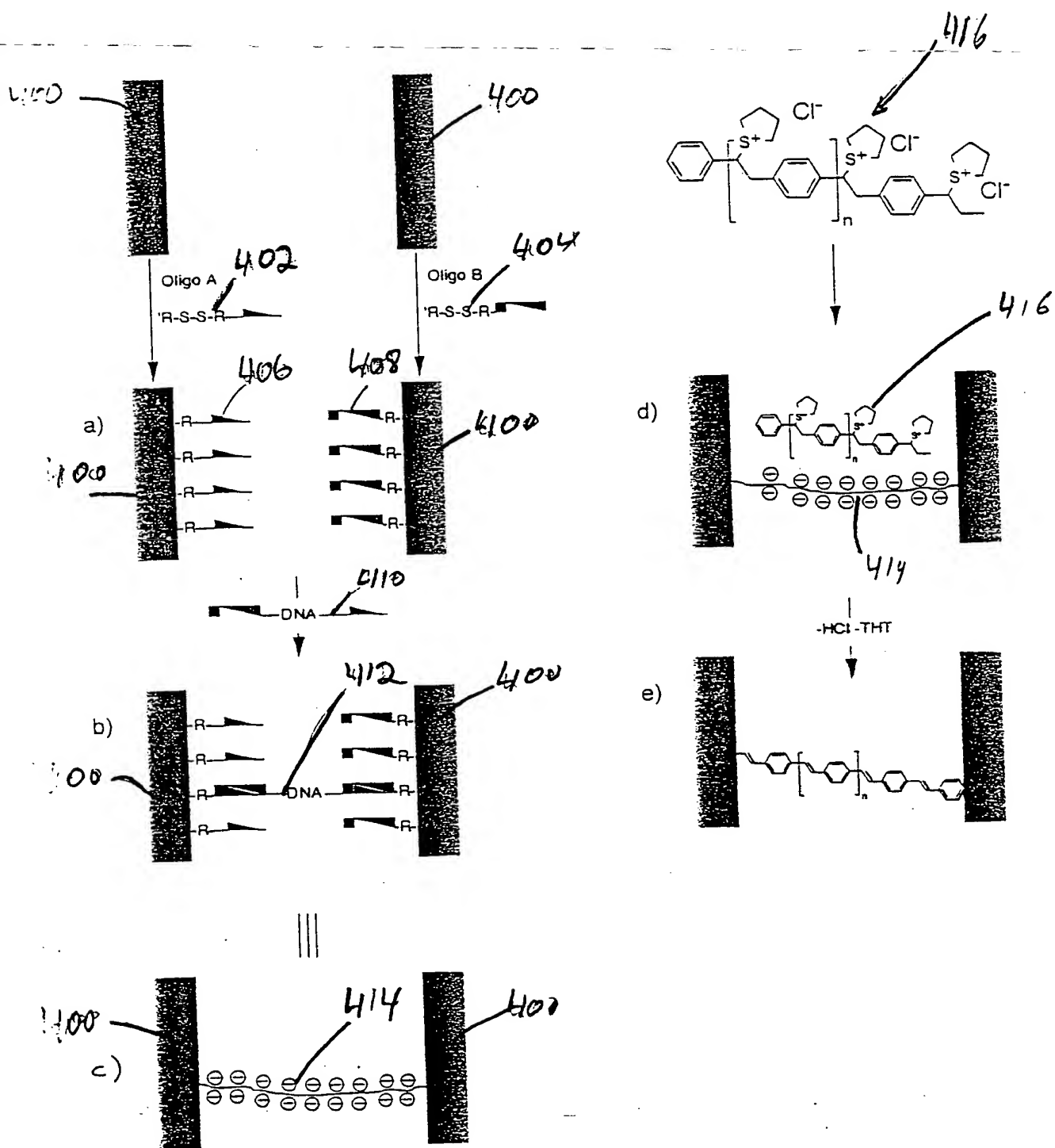


Fig. 4

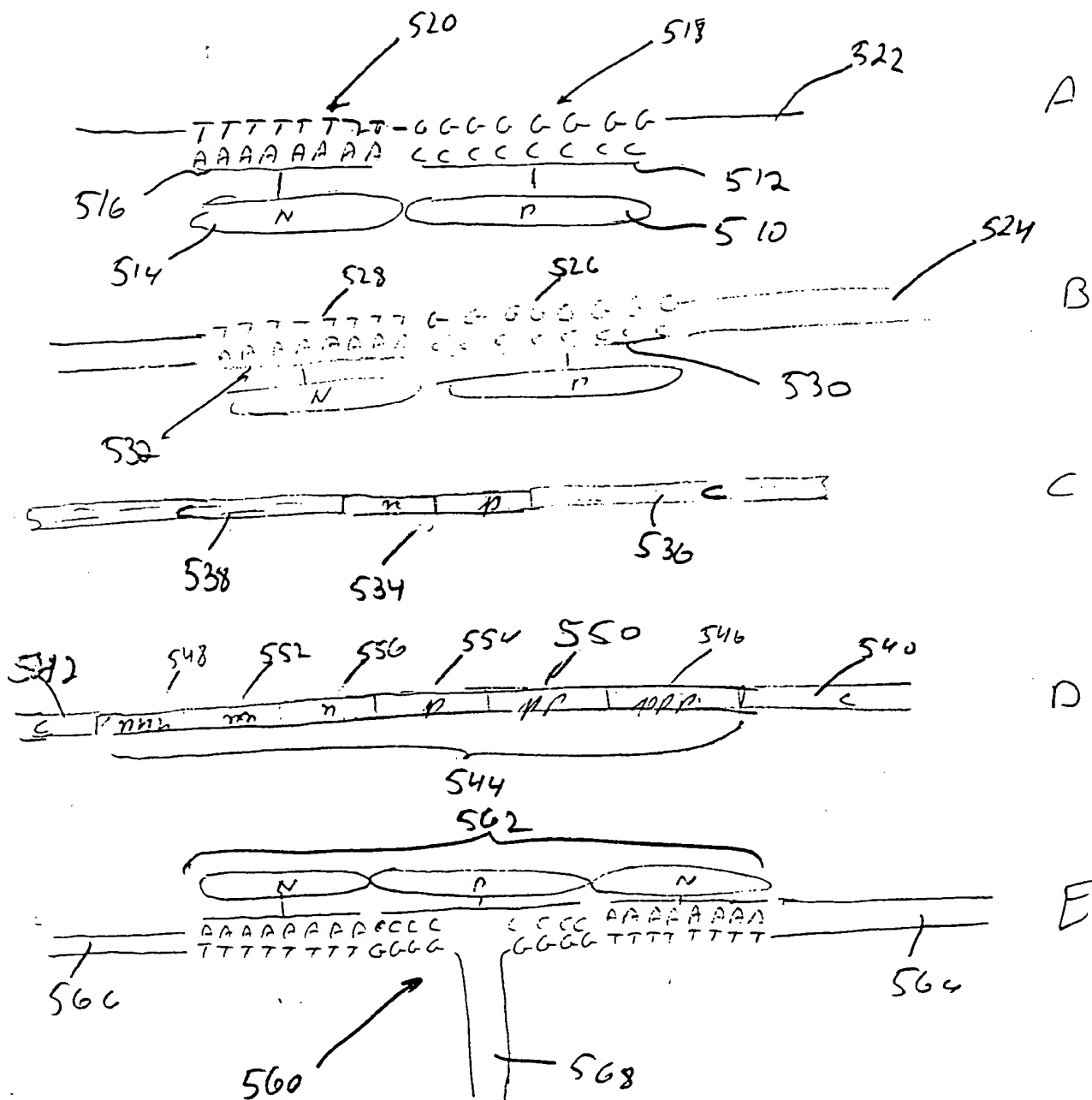
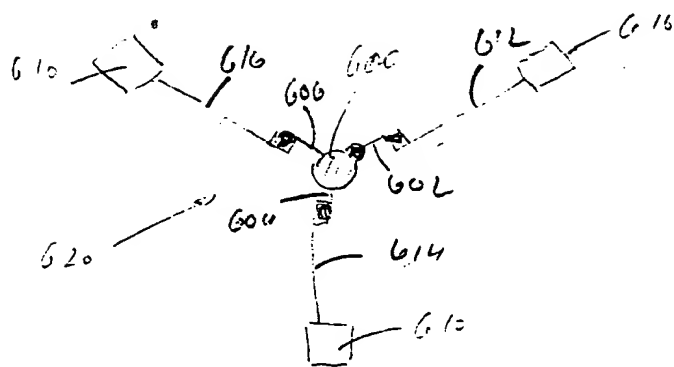
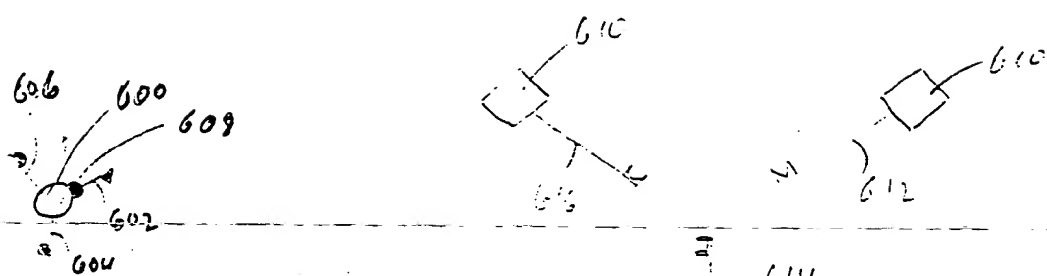


Fig. 5



Covalent binding  
Passivation

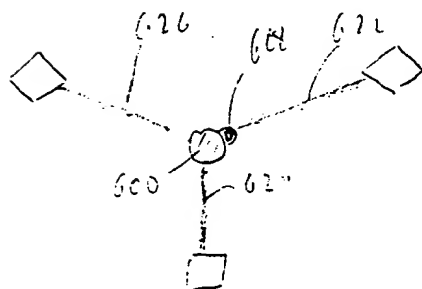
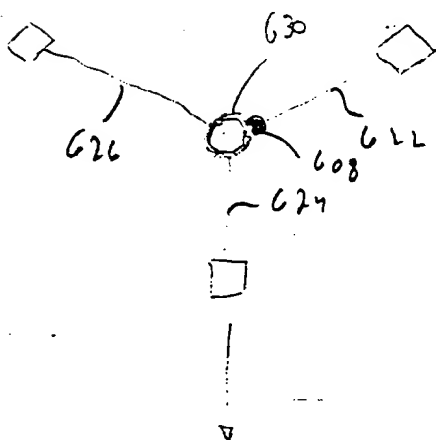


Fig. 6

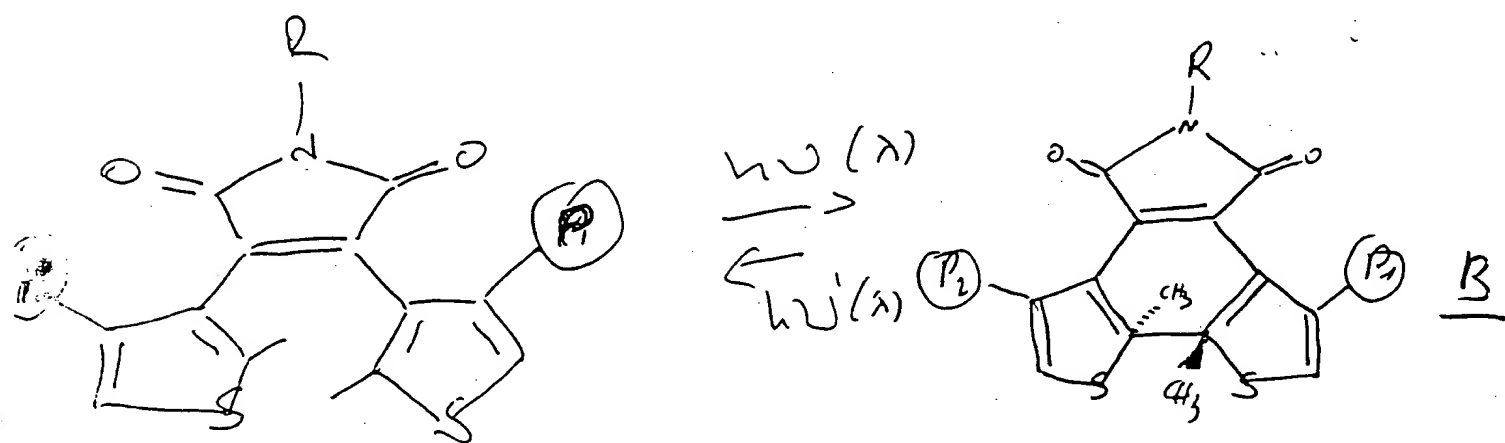
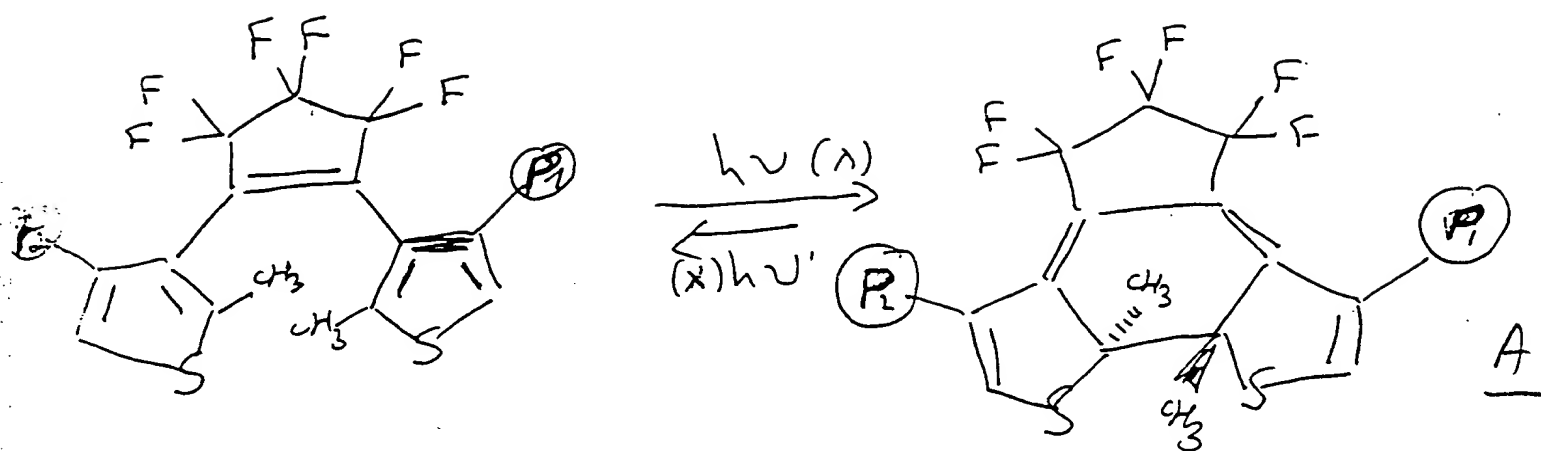


Fig 7

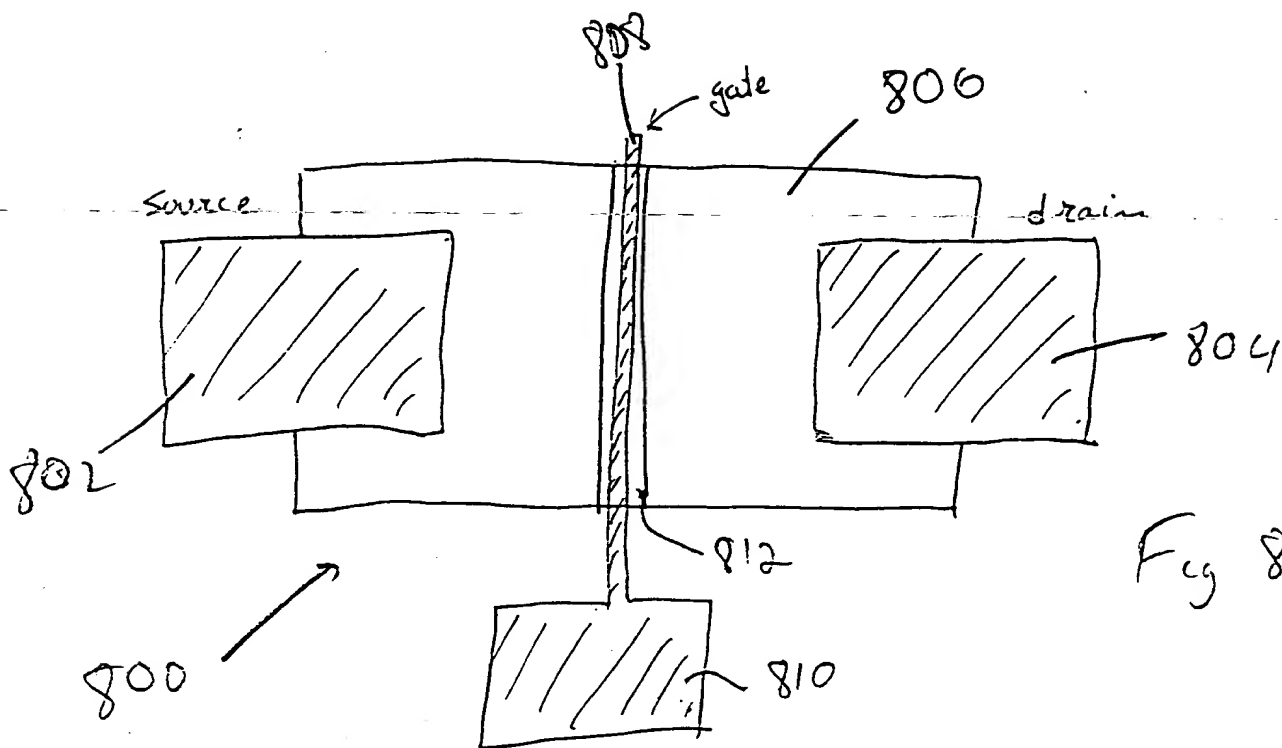


Fig 8A

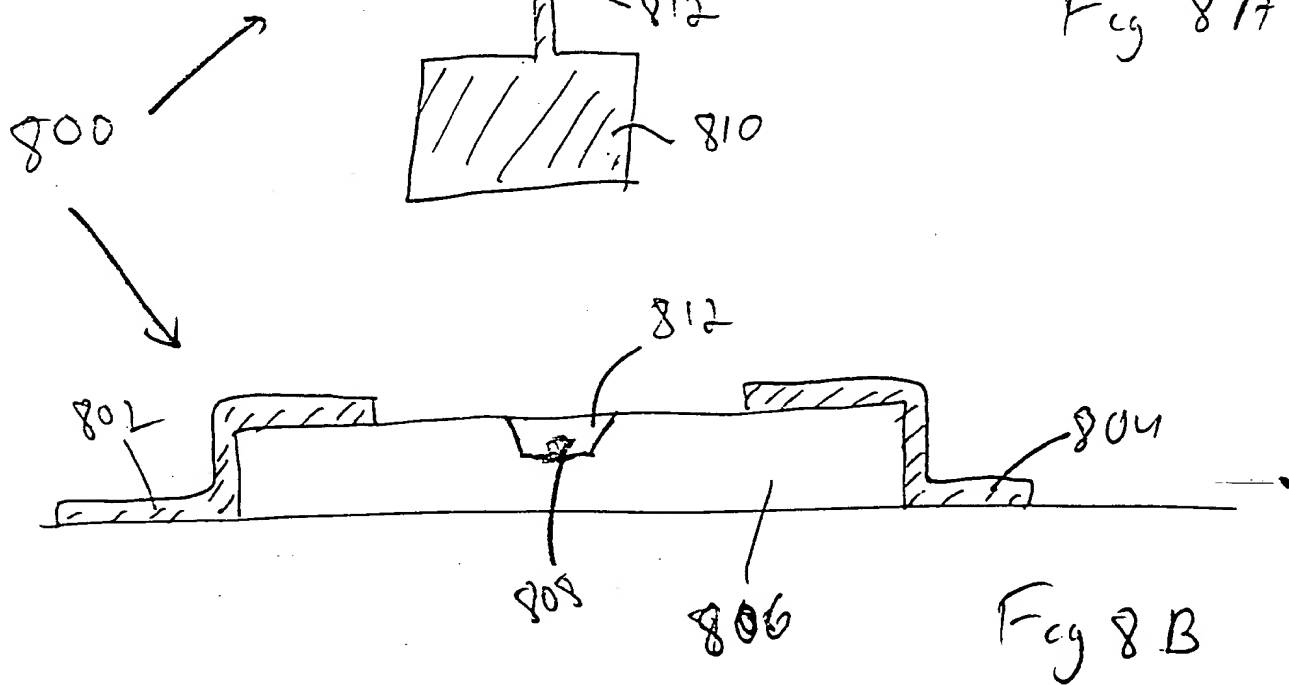


Fig 8B



Fig. 10

100.0 nm

50.0 nm

0.0 nm

1.50

1.00

0.50

0

1.50

nm

1.00

0.50

0

Fig 11A

60.0 nm

30.0 nm

0.0 nm

500

250

0

500

nm

250

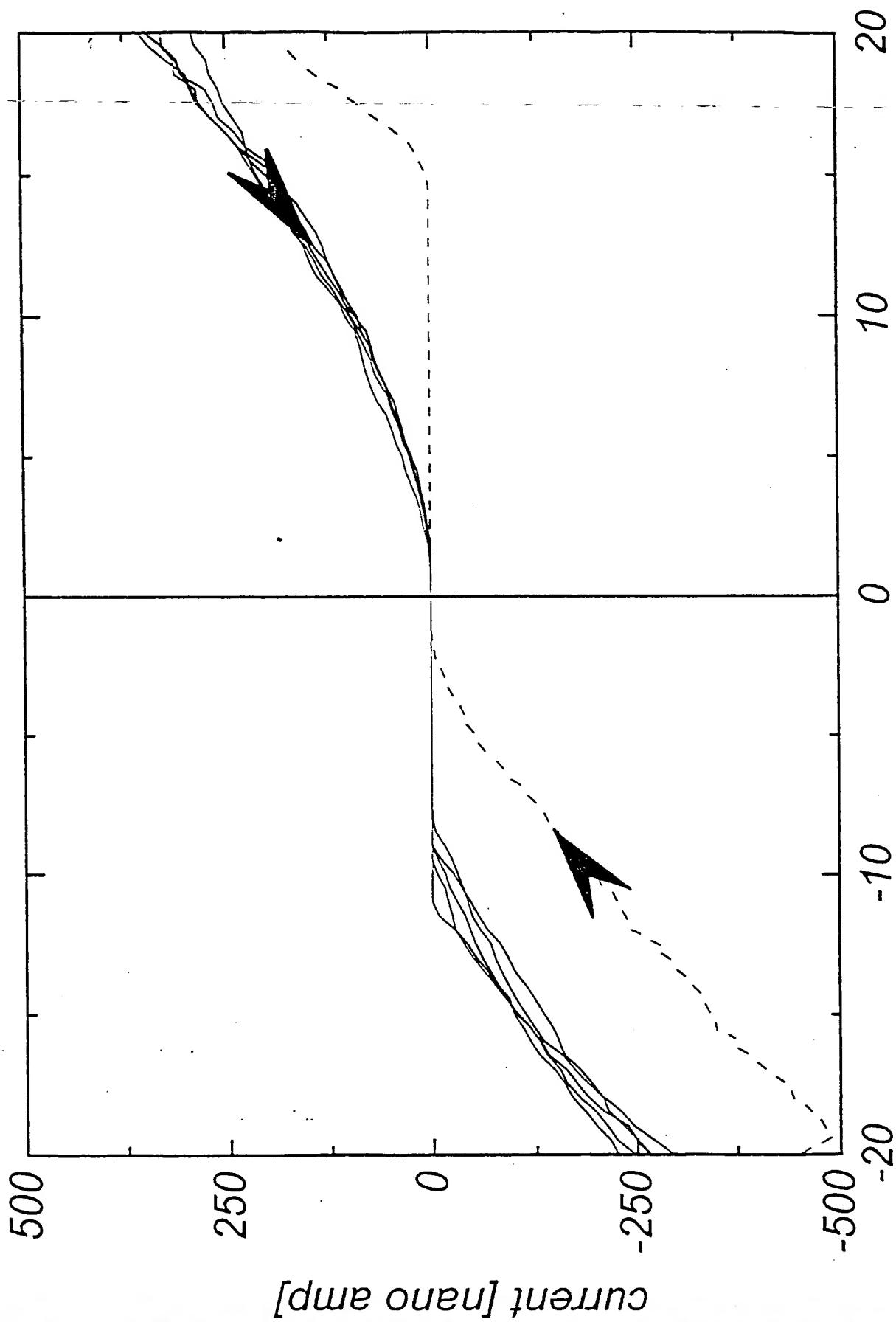
0

50

100

nm

Fig 11B



Bias [volt]  $f_{ig}$  12 A

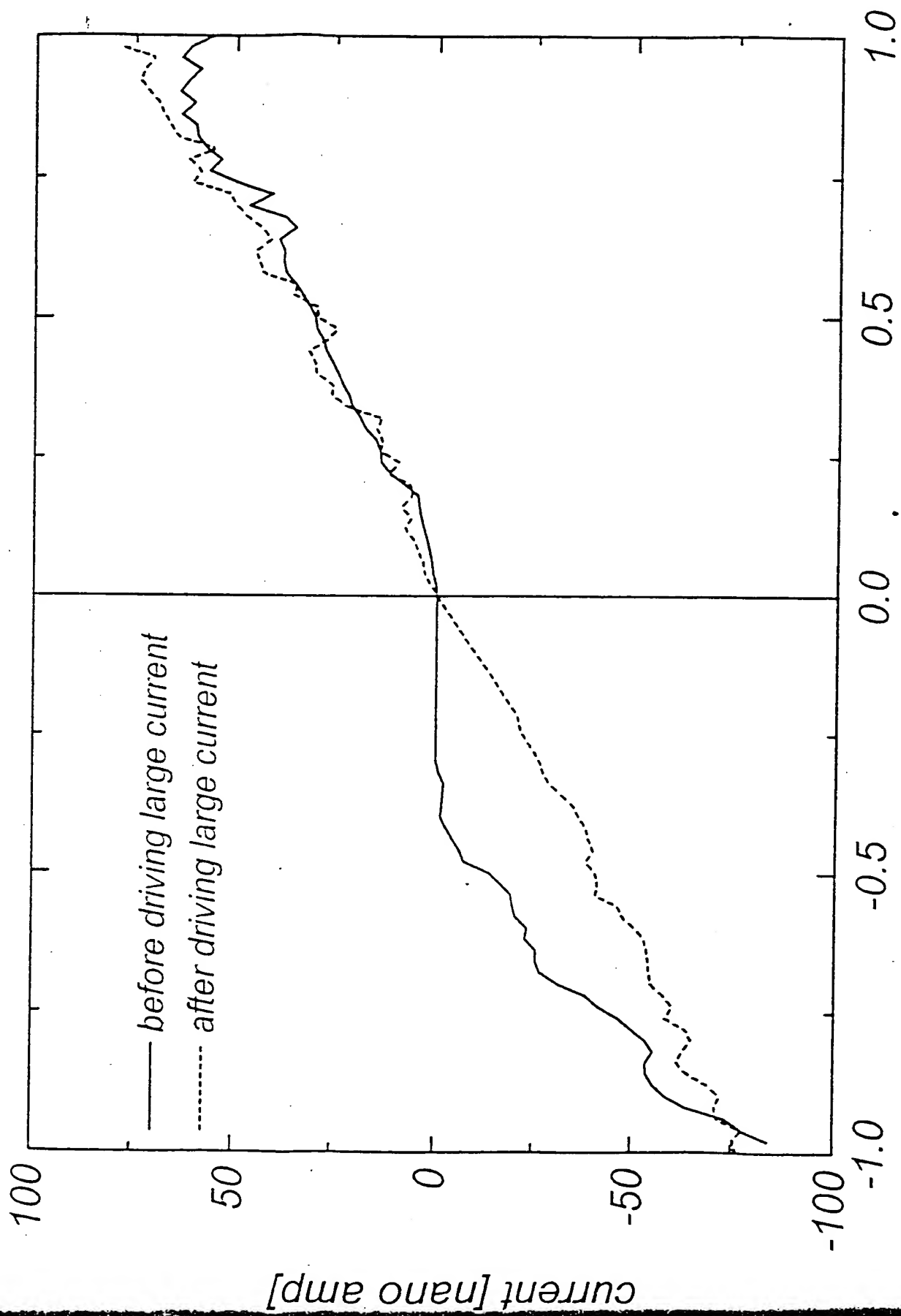


Fig 12B

Bias [volt]

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